



**Government of Khyber Pakhtunkhwa  
Health Department  
Directorate General Health Services  
Khyber Pakhtunkhwa Peshawar**

## **Standard Bidding Documents**

**For National Competitive Bidding  
Pakistan**

**For**

**SELECTION AND RATE CONTRACTING OF DRUGS / MEDICINES,  
MEDICAL DEVICES, SURGICAL DISPOSABLES &  
NON-DRUG ITEMS**

**FOR THE FINANCIAL YEARS 2021-23**

**MEDICINE COORDINATION CELL (MCC)**

**APRIL 2021**

## **PART ONE (UNCHANGEABLE)**

- Instructions to Bidders (ITB)
- General Conditions of Contract (GCC)

## Preface

These Bidding Documents have been prepared for use by procuring agencies and their implementing agencies in the procurement of goods through National Competitive Bidding (NCBs) as well International Competitive Bidding (ICBs) vide 41(g) KPP Rules 2014.

In order to simplify the preparation of bidding documents for each procurement, the Bidding Documents are grouped in two parts based on provisions which would remain the same for every procurement and that which are specific for each procurement. Provisions which are intended to be used unchanged are in Part one, which includes Section I, Instructions to Bidders, and Section II, General Conditions of Contract. Data and provisions specific to each procurement and contract are included in Part Two which is further organized into six sections. Sections I, II, III, IV, and V, respectively contain Invitation for Bids; Bid Data Sheet; Special Conditions of Contract; Schedule of Requirements; Technical Specifications; and the forms to be used, while Section VI is about Sample Forms.

This is Part one which is fixed and contains provisions which are to be used unchanged. Each section is prepared with notes intended only as information for the Procuring agency or the person drafting the bidding documents. They shall not be included in the final documents.

## Table of Contents - Part One

<b>PART ONE - SECTION I. INSTRUCTIONS TO BIDDERS</b>	<b>5</b>
Notes on the Instruction to Bidders	5
Table of Clauses	6
Instructions to Bidders	7-22
<b>PART ONE – SECTION II. GENERAL CONDITIONS OF CONTRACT</b>	<b>23</b>
Notes on the General Conditions of Contracts	24
Table of Clauses	25
General Conditions of Contracts	26-34

## Part One - Section I.

### Instructions to Bidders

#### **Notes on the Instructions to Bidders**

This section of the bidding documents provides the information necessary for bidders to prepare responsive bids, in accordance with the requirements of the Procuring agency. It also provides information on bid submission, opening, and evaluation, and on the award of contract.

Part One Section I contains provisions that are to be used unchanged. Part Two Section II (Bid Data Sheet) consists of provisions that supplement, amend, or specify in detail information or requirements included in Part One Section I and which are specific to each procurement.

Matters governing the performance of the Supplier, payments under the contract, or matters affecting the risks, rights, and obligations of the parties under the contract are not normally included in this section, but rather under Part one Section II, General Conditions of Contract, and/or Part Two Section III, Special Conditions of Contract. If duplication of a subject is inevitable in the other sections of the document prepared by the Procuring agency, care must be exercised to avoid contradictions between clauses dealing with the same matter.

These Instructions to Bidders will not be part of the contract.

## Table of Clauses

<b>A.</b>	<b>Introduction</b>	<b>7</b>
1.	Source of Funds	7
2.	Eligible Bidders	7
3.	Eligible Goods and Service	8
4.	Cost of Bidding	8
<b>B.</b>	<b>The Bidding Document</b>	<b>8</b>
5.	Content of Bidding Documents	8
6.	Clarification of Bidding Documents	8
7.	Amendment of Bidding Documents	9
<b>C.</b>	<b>Preparation of Bids</b>	<b>10</b>
8.	Language of Bid	10
9.	Documents Comprising the Bid	10
10.	Bid Form	10
11.	Bid Prices	10
12.	Bid Currencies	11
13.	Documents Establishing Bidder's Eligibility and Qualification	11
14.	Documents Establishing Goods' Eligibility and Conformity to Bidding Documents	11
15.	Bid Security	11
16.	Period of Validity of bids	12
17.	Format and Signing of Bid	12
<b>D.</b>	<b>Submission of Bids</b>	<b>12</b>
18.	Sealing and Marking of bids	12
19.	Deadline for Submission of bids	13
20.	Late bids	13
21.	Modification and Withdrawal of Bids	13
<b>E.</b>	<b>Opening and Evaluation of Bids</b>	<b>13</b>
22.	Opening of Bids by the Procuring Agency	13
23.	Clarification of Bids	13
24.	Preliminary Examination	14
25.	Evaluation and Comparison of Bids	14
26.	Contacting the Procuring Agency	19
<b>F.</b>	<b>Award of Contract</b>	<b>20</b>
27.	Post-Qualification	20
28.	Award Criteria	20
29.	Procuring Agency's Right to Vary Quantities at Time of Award	20
30.	Procuring Agency's Right to Accept Any Bid and To Reject Any or All Bids	20
31.	Notification of Award	20
32.	Signing of Contract	20
33.	Performance Security	22
34.	Corrupt or Fraudulent Practices	22
35.	Integrity Pact	22

## Instructions to Bidders

### A. Introduction

<b>1. Source of Funds</b>	1.1	The Procuring agency has received/applied for loan/grant/federal/provincial/local government funds from the source(s) indicated in the bidding data in various currencies towards the cost of the project /schemes specified in the bidding data and it is intended that part of the proceeds of this loan/grant/funds/ will be applied to eligible payments under the contract for which these bidding documents are issued.
	1.2	The funds referred to above in addition shall be “Public Fund” which according to 2 (l) of KPP Rules 2014 means ( i ) Provincial Consolidated Fund; ( ii) foreign assistance; ( iii) all moneys standing in the Public Account; and (iv) Funds of enterprises wholly or partly owned or managed or controlled by Government.
	1.3	Payment by the Fund will be made only at the request of the Procuring agency and upon approval by the Government of Khyber Pakhtunkhwa., and in case of a project will be subject in all respect to the terms and conditions of the agreement. The Project Agreement prohibits a withdrawal from the allocated fund account for the purpose of any payment to persons or entities, or for any import of goods, if such payment or import, to the knowledge of the Federal Government/ Khyber Pakhtunkhwa Government, is prohibited by a decision of the United Nations Security Council taken under Chapter VII of the Charter of the United Nations. No party other than the Procuring agency shall derive any rights from the Project Agreement or have any claim to the allocated fund proceeds.
<b>2. Eligible Bidders</b>	2.1	This Invitation for Bids is open to all suppliers from eligible source as defined in the KPP Rules, 2014 and its Bidding Documents except as provided hereinafter.
	2.2	Bidders should not be associated, or have been associated in the past, directly or indirectly, with a firm or any of its affiliates which have been engaged by the Procuring agency to provide consulting services for the preparation of the design, specifications, and other documents to be used for the procurement of the goods to be purchased under this Invitation for Bids.
	2.3	Government-owned enterprises in the Province of Khyber Pakhtunkhwa may participate only if they are legally and financially autonomous, if they operate under commercial law, and if they are not a dependent agency of the Government of Khyber Pakhtunkhwa.
	2.4	Bidders shall not be eligible to bid if they are under a declaration of ineligibility for corrupt and fraudulent practices issued by any government organization in accordance with the Section 44(1) KPP Rules 2014.

<b>3. Eligible Goods and Services</b>	3.1	All goods and related services to be supplied under the contract shall have their origin in eligible source countries of the world with whom the Islamic Republic of Pakistan has commercial relations and its Bidding Documents and all expenditures made under the contract will be limited to such goods and services.
	3.2	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. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized product results that is substantially different in basic characteristics or in purpose or utility from its components.
	3.3	The origin of goods and services is distinct from the nationality of the Bidder.
<b>4. Cost of Bidding</b>	4.1	The Bidder shall bear all costs associated with the preparation and submission of its bid, and the Procuring agency named in the Bid Data Sheet, hereinafter referred to as “the Procuring agency,” will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.
		<b>B. The Bidding Documents</b>
<b>5. Content of Bidding Documents</b>	5.1	The bidding documents include: <ul style="list-style-type: none"> <li>a) Instructions to Bidders (ITB)</li> <li>b) Bid Data Sheet</li> <li>c) General Conditions of Contract (GCC)</li> <li>d) Special Conditions of Contract (SCC)</li> <li>e) Schedule of Requirements</li> <li>f) Technical Specifications</li> <li>g) Bid Form and Price Schedules</li> <li>h) Bid Security Form</li> <li>i) Contract Form</li> <li>j) Performance Security Form</li> <li>k) Manufacturer’s Authorization Form</li> </ul>
	5.2	The Bidder is expected to examine all instructions, forms, terms, and specifications in the bidding documents. Failure to furnish all information required by the bidding documents or to submit a bid not substantially responsive to the bidding documents in every respect will be at the Bidder’s risk and may result in the rejection of its bid.
<b>6. Clarification of Bidding Documents</b>	6.1	An interested Bidder requiring any clarification of the bidding documents may notify the Procuring agency in writing. The Bidding Procuring agency will respond in writing to any request for Document’s clarification of the bidding documents which it receives no later than three working days prior to the deadline for the submission of bids prescribed in the Bid Data Sheet. Written copies of the Procuring agency’s response (including an explanation of the query but without identifying the source of inquiry) will be sent to all interested bidders that have received the bidding documents.

<b>7. Amendment of Bidding Documents</b>	7.1	At any time prior to the deadline for submission of bids, the Procuring agency, for any reason, whether at its own initiative or in response to a clarification requested by a interested Bidder, may modify the bidding documents by amendment.
	7.2	All interested bidders that have received the bidding documents will be notified of the amendment in writing and will be binding on them.
	7.3	In order to allow interested bidders reasonable time in which to take the amendment into account in preparing their bids, the Procuring agency, at its discretion, may extend the deadline for the submission of bids.
<b>C. Preparation of Bids</b>		
<b>8. Language of Bid</b>	8.1	The bid prepared by the Bidder, as well as all correspondence and documents relating to the bid exchanged by the Bidder and the Procuring agency shall be written in the language specified in the Bid Data Sheet. Supporting documents and printed literature furnished by the Bidder may be in another language provided they are accompanied by an accurate translation of the relevant passages in the language specified in the Bid Data Sheet, in which case, for purposes of interpretation of the Bid, the translation shall govern.
<b>9. Documents Comprising the Bid</b>	9.1	The bid prepared by the Bidder shall comprise the following components: <ul style="list-style-type: none"> <li>a) A Bid Form and a Price Schedule completed in accordance with ITB Clauses 10, 11, and 12</li> <li>b) documentary evidence established in accordance with ITB Clause 13 that the Bidder is eligible to bid and is qualified to perform the contract if its bid is accepted;</li> <li>c) documentary evidence established in accordance with ITB Clause 14 that the goods and ancillary services to be supplied by the Bidder are eligible goods and services and conform to the bidding documents; and</li> </ul> Bid security furnished in accordance with ITB Clause 15.
<b>10. Bid Form</b>	10.1	The Bidder shall complete the Bid Form and the appropriate Price Schedule furnished in the bidding documents, indicating the goods to be supplied, a brief description of the goods, and their country of origin, quantity, and prices.
<b>11. Bid Prices</b>	11.1	The Bidder shall indicate on the appropriate Price Schedule the unit prices (where applicable) and total bid price of the goods it proposes to supply under the contract.
	11.2	Prices indicated on the Price Schedule shall be delivered duty paid (DDP) prices. The price of other (incidental) services, if any, listed in the Bid Data Sheet will be entered separately.
	11.3	The Bidder's separation of price components in accordance with ITB Clause 11.2 above will be solely for the purpose of facilitating the comparison of bids by the Procuring agency and will not in any way limit the Procuring agency's right to contract on any of the terms offered.
	11.4	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 Bid Data Sheet. A bid submitted with an adjustable price quotation will be treated as nonresponsive and will be rejected, pursuant to ITB Clause 24. If, however, in accordance with the Bid Data Sheet, prices quoted by the Bidder shall be subject to adjustment during the performance of the contract, a bid submitted with a fixed price quotation will not be rejected, but the price adjustment would be treated as zero.

<b>12. Bid Currencies</b>	12.1	Prices shall be quoted in Pak Rupees unless otherwise specified in the Bid Data Sheet.
<b>13. Documents Establishing Bidder's</b>	13.1	Pursuant to ITB Clause 9, the Bidder shall furnish, as part of its bid, documents establishing the Bidder's eligibility to bid and its qualifications to perform the contract if its bid is accepted.
<b>Eligibility and Qualification</b>	13.2	The documentary evidence of the Bidder's eligibility to bid shall establish to the Procuring agency's satisfaction that the Bidder, at the time of submission of its bid, is from an eligible country as defined under ITB Clause 3.
	13.3	<p>The documentary evidence of the Bidder's qualifications to perform the contract if its bid is accepted shall establish to the Procuring agency's satisfaction:</p> <ul style="list-style-type: none"> <li>a) that, in the case of a Bidder offering to supply goods under the contract which the Bidder did not manufacture or otherwise produce, the Bidder has been duly authorized by the goods' Manufacturer or producer to supply the goods in the Procuring agency's country;</li> <li>b) that the Bidder has the financial, technical, and production capability necessary to perform the contract;</li> <li>c) that, in the case of a Bidder not doing business within the Procuring agency's country, the Bidder is or will be (if awarded the contract) represented by an Agent in that country 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; and</li> <li>d) That the Bidder meets the qualification criteria listed in the Bid Data Sheet.</li> </ul>
<b>14. Documents Establishing Goods' Eligibility and Conformity to Bidding Documents</b>	14.1	Pursuant to ITB Clause 9, the Bidder shall furnish, as part of its bid, documents establishing the eligibility and conformity to the bidding documents of all goods and services which the Bidder proposes to supply under the contract.
	14.2	The documentary evidence of the eligibility of the goods and services shall consist of a statement in the Price Schedule of the country of origin of the goods and services offered which shall be confirmed by a certificate of origin issued at the time of shipment.

	14.3	<p>The documentary evidence of conformity of the goods and services to the bidding documents may be in the form of literature, drawings, and data, and shall consist of:</p> <ul style="list-style-type: none"> <li>a) a detailed description of the essential technical and performance characteristics of the goods;</li> <li>b) a list giving full particulars, including available sources and current prices of spare parts, special tools, etc., necessary for the proper and continuing functioning of the goods for a period to be specified in the Bid Data Sheet, following commencement of the use of the goods by the Procuring agency; and</li> </ul>
		c) an item-by-item commentary on the Procuring agency's Technical Specifications demonstrating substantial responsiveness of the goods and services to those specifications, or a statement of deviations and exceptions to the provisions of the Technical Specifications.
	14.4	For purposes of the commentary to be furnished pursuant to ITB Clause 14.3(c) above, the Bidder shall note that standards for workmanship, material, and equipment, as well as references to brand names or catalogue numbers designated by the Procuring agency in its Technical Specifications, are intended to be descriptive only and not restrictive. The Bidder may substitute alternative standards, brand names, and/or catalogue numbers in its bid, provided that it demonstrates to the Procuring agency's satisfaction that the substitutions ensure substantial equivalence to those designated in the Technical Specifications.
<b>15. Bid Security</b>	15.1	Pursuant to ITB Clause 9, the Bidder shall furnish, as part of its bid, a bid security in the amount specified in the Bid Data Sheet.
	15.2	The bid security is required to protect the Procuring agency against the risk of Bidder's conduct which would warrant the security's forfeiture, pursuant to ITB Clause 15.7.
	15.3	<p>The bid security shall be in Pak. Rupees and shall be in one of the following forms:</p> <ul style="list-style-type: none"> <li>a) a bank guarantee or an irrevocable letter of credit issued by a reputable bank located in the Procuring agency's country, in the form provided in the bidding documents or another form acceptable to the Procuring agency and valid for thirty (30) days beyond the validity of the bid; or</li> </ul> <p>Irrevocable encashable on-demand Bank call-deposit.</p>
	15.4	Any bid not secured in accordance with ITB Clauses 15.1 and 15.3 will be rejected by the Procuring agency as non-responsive, pursuant to ITB Clause 24.
	15.5	Unsuccessful bidders' bid security will be discharged or returned as promptly as possible but not later than thirty (30) days after the expiration of the period of bid validity prescribed by the Procuring agency pursuant to ITB Clause 16.
	15.6	The successful Bidder's bid security will be discharged upon the Bidder signing the contract, pursuant to ITB Clause 32, and furnishing the performance security, pursuant to ITB Clause 33.

	15.7	<p>The bid security may be forfeited:</p> <ul style="list-style-type: none"> <li>a) if a Bidder withdraws its bid during the period of bid validity specified by the Bidder on the Bid Form; or</li> <li>b) in the case of a successful Bidder, if the Bidder fails: <ul style="list-style-type: none"> <li>i. to sign the contract in accordance with ITB Clause 32;</li> <li>or</li> </ul> </li> </ul> <p>to furnish performance security in accordance with ITB Clause 33.</p>
<b>16. Period of Validity of Bids</b>	16.1	Bids shall remain valid for the period specified in the Bid Data Sheet after the date of bid opening prescribed by the Procuring agency, pursuant to ITB Clause 19. A bid valid for a shorter period shall be rejected by the Procuring agency as non-responsive.
	16.2	In exceptional circumstances, the Procuring agency may solicit the Bidder's consent to an extension of the period of validity. The request and the responses thereto shall be made in writing. The bid security provided under ITB Clause 15 shall also be suitably extended. A Bidder may refuse the request without forfeiting its bid security. A Bidder granting the request will not be required nor permitted to modify its bid, except as provided in the bidding document.
<b>17. Format and Signing of Bid</b>	17.1	The Bidder shall prepare an original and the number of copies of the bid indicated in the Bid Data Sheet, clearly marking each "ORIGINAL BID" and "COPY OF BID," as appropriate. In the event of any discrepancy between them, the original shall govern.
	17.2	The original and the copy or copies of the bid shall be typed or written in indelible ink and shall be signed by the Bidder or a person or persons duly authorized to bind the Bidder to the contract. All pages of the bid, except for un-amended printed literature, shall be initialed by the person or persons signing the bid.
	17.3	Any interlineations, erasures, or overwriting shall be valid only if they are initialed by the person or persons signing the bid.
	17.4	The Bidder shall furnish information as described in the Form of Bid on commissions or gratuities, if any, paid or to be paid to agents relating to this Bid, and to contract execution if the Bidder is awarded the contract.
		<b>D. Submission of Bids</b>
<b>18. Sealing and Marking of Bids</b>	18.1	The Bidder shall seal the original and each copy of the bid in separate envelopes, duly marking the envelopes as "ORIGINAL" and "COPY." The envelopes shall then be sealed in an outer envelope.
	18.2	<p>The inner and outer envelopes shall:</p> <ul style="list-style-type: none"> <li>a. be addressed to the Procuring agency at the address given in the Bid Data Sheet; and</li> </ul> <p>bear the Project name indicated in the Bid Data Sheet, the Invitation for Bids (IFB) title and number indicated in the Bid Data Sheet, and a statement: "DO NOT OPEN BEFORE," to be completed with the time and the date specified in the Bid Data Sheet, pursuant to ITB Clause 2.2.</p>
	18.3	The inner envelopes shall also indicate the name and address of the Bidder to enable the bid to be returned unopened in case it is declared "late".
	18.4	If the outer envelope is not sealed and marked as required by ITB Clause 18.2, the Procuring agency will assume no responsibility for the bid's misplacement or premature opening.

<b>19. Deadline for Submission of Bids</b>	19.1	Bids must be received by the Procuring agency at the address specified under ITB Clause 18.2 no later than the time and date specified in the Bid data sheet.
	19.2	The Procuring agency may, at its discretion, extend this deadline for the submission of bids by amending the bidding documents in accordance with ITB Clause 7, in which case all rights and obligations of the Procuring agency and bidders previously subject to the deadline will thereafter be subject to the deadline as extended.
<b>20. Late Bids</b>	20.1	Any bid received by the Procuring agency after the deadline for submission of bids prescribed by the Procuring agency pursuant to ITB Clause 19 will be rejected and returned unopened to the Bidder.
<b>21. Modification And Withdrawal of Bids</b>	21.1	The Bidder may modify or withdraw its bid after the bid's submission, provided that written notice of the modification, including substitution or withdrawal of the bids, is received by the Procuring agency prior to the deadline prescribed for submission of bids.
	21.2	The Bidder's modification or withdrawal notice shall be prepared, sealed, marked, and dispatched in accordance with the provisions of ITB Clause 18. by a signed confirmation copy, postmarked not later than the deadline for submission of bids.
	21.3	No bid may be modified after the deadline for submission of bids.
	21.4	No bid may be withdrawn in the interval between the deadline for submission of bids and the expiration of the period of bid validity specified by the Bidder on the Bid Form. Withdrawal of a bid during this interval may result in the Bidder's forfeiture of its bid security, pursuant to the ITB Clause 15.7.
		<b>E. Opening and Evaluation of Bids</b>
<b>22. Opening of Bids by the Procuring Agency</b>	22.1	The Procuring agency will open all bids in the presence of bidders' representatives who choose to attend, at the time, on the date, and at the place specified in the Bid Data Sheet. The bidders' representatives who are present shall sign a register evidencing their attendance.
	22.2	The bidders' names, bid modifications or withdrawals, bid prices, discounts, and the presence or absence of requisite bid security and such other details as the Procuring agency, at its discretion, may consider appropriate, will be announced at the opening. No bid shall be rejected at bid opening, except for late bids, which shall be returned unopened to the Bidder pursuant to ITB Clause 20.
	22.3	Bids (and modifications sent pursuant to ITB Clause 21.2) that are not opened and read out at bid opening shall not be considered further for evaluation, irrespective of the circumstances. Withdrawn bids will be returned unopened to the bidders.
	22.4	The Procuring agency will prepare minutes of the bid opening.
<b>23. Clarification of Bids</b>	23.1	During evaluation of the bids, the Procuring agency may, at its discretion, ask the Bidder for a clarification of its bid. The Bids request for clarification and the response shall be in writing, and no change in the prices or substance of the bid shall be sought, offered, or permitted.

<b>24. Preliminary Examination</b>	24.1	The Procuring agency will examine the bids to determine whether they are complete, whether any computational errors have been made, whether required sureties have been furnished, whether the documents have been properly signed, and whether the bids are generally in order.
	24.2	Arithmetical errors will be rectified on the following basis. 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. If the Supplier does not accept the correction of the errors, its bid will be rejected, and its bid security may be forfeited. If there is a discrepancy between words and figures, the amount in words will prevail.
	24.3	The Procuring agency may waive any minor informality, nonconformity, or irregularity in a bid which does not constitute a material deviation, provided such waiver does not prejudice or affect the relative ranking of any Bidder.
	24.4	Prior to the detailed evaluation, pursuant to ITB Clause 25 the Procuring agency will determine the substantial responsiveness of each bid to the bidding documents. For purposes of these Clauses, a substantially responsive bid is one which conforms to all the terms and conditions of the bidding documents without material deviations. Deviations from, or objections or reservations to critical provisions, such as those concerning Bid Security (ITB Clause 15), Applicable Law (GCC Clause 30), and Taxes and Duties (GCC Clause 32), will be deemed to be a material deviation. The Procuring agency's determination of a bid's responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence.
	24.5	If a bid is not substantially responsive, it will be rejected by the Procuring agency and may not subsequently be made responsive by the Bidder by correction of the nonconformity.
<b>25. Evaluation and Comparison of Bids</b>	25.1	The Procuring agency will evaluate and compare the bids which have been determined to be substantially responsive, pursuant to ITB Clause 24.
	25.2	The Procuring agency's evaluation of a bid will be on delivered duty paid (DDP) price inclusive of prevailing duties and will exclude any allowance for price adjustment during the period of execution of the contract, if provided in the bid.
	25.3	<p>The Procuring agency's evaluation of a bid will take into account, in addition to the bid price quoted in accordance with ITB Clause 11.2, one or more of the following factors as specified in the Bid Data Sheet, and quantified in ITB Clause 25.4:</p> <ul style="list-style-type: none"> <li>a. incidental costs</li> <li>b. delivery schedule offered in the bid;</li> <li>c. deviations in payment schedule from that specified in the Special Conditions of Contract;</li> <li>d. the cost of components, mandatory spare parts, and service;</li> <li>e. the availability Procuring agency of spare parts and after-sales services for the equipment offered in the bid;</li> <li>f. the projected operating and maintenance costs during the life of the equipment; the performance and productivity of the</li> </ul>

		<p>equipment offered; and/or</p> <p>g. other specific criteria indicated in the Bid Data Sheet and/or</p> <p>h. in the Technical Specifications.</p>
--	--	--

	25.4	<p>For factors retained in the Bid Data Sheet pursuant to ITB 25.3, one or more of the following quantification methods will be applied, as detailed in the Bid Data Sheet:</p> <ul style="list-style-type: none"> <li>a. Incidental costs provided by the bidder will be added by Procuring agency to the delivered duty paid (DDP) price at the final destination.</li> <li>b. Delivery schedule. <ul style="list-style-type: none"> <li>i. The Procuring agency requires that the goods under the Invitation for Bids shall be delivered at the time specified in the Schedule of Requirements which will be treated as the base, a delivery “adjustment” will be calculated for bids by applying a percentage, specified in the Bid Data Sheet, of the DDP price for each week of delay beyond the base, and this will be added to the bid price for evaluation. No credit shall be given to early delivery.</li> <li>or</li> <li>ii. The goods covered under this invitation are required to be delivered (shipped) within an acceptable range of weeks specified in the Schedule of Requirement. No credit will be given to earlier deliveries, and bids offering delivery beyond this range will be treated as non-responsive. Within this acceptable range, an adjustment per week, as specified in the Bid Data Sheet, will be added for evaluation to the bid price of bids offering deliveries later than the earliest delivery period specified in the Schedule of Requirements.</li> <li>or</li> <li>iii. The goods covered under this invitation are required to be delivered in partial shipments, as specified in the Schedule of Requirements. Bids offering deliveries earlier or later than the specified deliveries will be adjusted in the evaluation by adding to the bid price a factor equal to a percentage, specified in the Bid Data Sheet, of DDP price per week of variation from the specified delivery schedule.</li> </ul> </li> <li>c. Deviation in payment schedule: <ul style="list-style-type: none"> <li>i. Bidders shall state their bid price for the payment schedule outlined in the SCC. Bids will be evaluated on the basis of this base price. Bidders are, however, permitted to state an alternative payment schedule and indicate the reduction in bid price they wish to offer for such alternative payment schedule. The Procuring agency may consider the alternative payment schedule offered by the selected Bidder.</li> <li>or</li> <li>ii. The SCC stipulates the payment schedule offered by the Procuring agency. If a bid deviates from the schedule and if such deviation is considered acceptable to the Procuring agency, the bid will be evaluated by calculating interest earned for any earlier payments involved in the terms outlined in the bid as compared with those stipulated in this</li> </ul> </li> </ul>
--	------	---

		<p>invitation, at the rate per annum specified in the Bid Data Sheet.</p> <p>d. Cost of spare parts.</p> <p>i. The list of items and quantities of major assemblies, components, and selected spare parts, likely to be required during the initial period of operation specified in the Bid Data Sheet, is annexed to the Technical Specifications. The total cost of these items, at the unit prices quoted in each bid, will be added to the bid price.</p> <p>or</p> <p>ii. The Procuring agency will draw up a list of high- usage and high-value items of components and spare parts, along with estimated quantities of usage in the initial period of operation specified in the Bid Data Sheet. The total cost of these items and quantities will be computed from spare parts unit prices submitted by the Bidder and added to the bid price.</p> <p>or</p> <p>iii. The Procuring agency will estimate the cost of spare parts usage in the initial period of operation specified in the Bid Data Sheet, based on information furnished by each Bidder, as well as on past experience of the Procuring agency or other procuring agencies in similar situations. Such costs shall be added to the bid price for evaluation.</p> <p>e. Spare parts and after sales service facilities in the Procuring agency's country.</p> <p>The cost to the Procuring agency of establishing the minimum service facilities and parts inventories, as outlined in the Bid Data Sheet or elsewhere in the bidding documents, if quoted separately, shall be added to the bid price.</p> <p>f. Operating and maintenance costs.</p> <p>Since the operating and maintenance costs of the goods under procurement form a major part of the life cycle cost of the equipment, these costs will be evaluated in accordance with the criteria specified in the Bid Data Sheet or in the Technical Specifications.</p> <p>g. Performance and productivity of the equipment.</p> <p>i. Bidders shall state the guaranteed performance or efficiency in response to the Technical Specification. For each drop in the performance or efficiency below the norm of 100, an adjustment for an amount specified in the Bid Data Sheet will be added to the bid price, representing the capitalized cost of additional operating costs over the life of the plant, using the methodology specified in the Bid Data Sheet or in the Technical Specifications.</p> <p>or</p> <p>ii. Goods offered shall have a minimum productivity specified under the relevant provision in the Technical</p>
--	--	--

		<p>Specifications to be considered responsive. Evaluation shall be based on the cost per unit of the actual productivity of goods offered in the bid, and adjustment will be added to the bid price using the methodology specified in the Bid Data Sheet or in the Technical Specifications.</p> <p>h. Specific additional criteria indicated in the Bid Data Sheet and/or in the Technical Specifications.</p> <p>The relevant evaluation method shall be detailed in the Bid Data Sheet and/or in the Technical Specifications</p>
--	--	---

<b>Alternative</b>	25.4	25.4 Merit Point System:	
		The following merit point system for weighing evaluation factors can be applied if none of the evaluation methods listed in 25.4 above has been retained in the Bid Data Sheet. The number of points allocated to each factor shall be specified in the Bid Data Sheet.	
		[In the Bid Data Sheet, choose from the range of]	
		Evaluated price of the goods	60 to 90
		Cost of common list spare parts	0 to 20
		Technical features, and maintenance and operating costs	0 to 20
		Availability of service and spare parts	0 to 20
		Standardization	0 to 20
		Total	100
		The bid scoring the highest number of points will be deemed to be the lowest evaluated bid.	
<b>26. Contacting the Procuring Agency</b>	26.1	Subject to ITB Clause 23, no Bidder shall contact the Procuring agency on any matter relating to its bid, from the time of the bid opening to the time the contract is awarded. If the Bidder wishes to bring additional information to the notice of the Procuring agency, it should do so in writing.	
	26.2	Any effort by a Bidder to influence the Procuring agency in its decisions on bid evaluation, bid comparison, or contract award may result in the rejection of the Bidder's bid.	

		<b>F. Award of Contract</b>
<b>27. Post- qualification</b>	27.1	In the absence of prequalification, the Procuring agency will determine to its satisfaction whether the Bidder that is selected as having submitted the lowest evaluated responsive bid is qualified to perform the contract satisfactorily, in accordance with the criteria listed in ITB Clause 13.3.
	27.2	The determination will take into account the Bidder's financial, technical, and production capabilities. It will be based upon an examination of the documentary evidence of the Bidder's qualifications submitted by the Bidder, pursuant to ITB Clause 13.3, as well as such other information as the Procuring agency deems necessary and appropriate.
	27.3	An affirmative determination will be a prerequisite for award of the contract to the Bidder. A negative determination will result in rejection of the Bidder's bid, in which event the Procuring agency will proceed to the next lowest evaluated bid to make a similar determination of that Bidder's capabilities to perform satisfactorily.
<b>28. Award Criteria</b>	28.1	Subject to ITB Clause 30, the Procuring agency will award the contract to the successful Bidder whose bid has been determined to be substantially responsive and has been determined to be the lowest evaluated bid, provided further that the Bidder is determined to be qualified to perform the contract satisfactorily.
<b>29. Procuring agency's Right to Vary Quantities at Time of Award</b>	29.1	The Procuring agency reserves the right at the time of contract award to increase or decrease, by the percentage indicated in the Bid Data Sheet, the quantity of goods and services originally specified in the Schedule of Requirements without any change in unit price or other terms and conditions.
<b>30. Procuring agency's Right to Accept any Bid and to Reject any or All Bids</b>	30.1	The Procuring agency 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 affected Bidder or bidders or any obligation to inform the affected Bidder or bidders of the grounds for the Procuring agency's action.
<b>31. Notification of Award</b>	31.1	Prior to the expiration of the period of bid validity, the Procuring agency will notify the successful Bidder in writing by registered letter or by cable, to be confirmed in writing by registered letter, that its bid has been accepted.
	31.2	The notification of award will constitute the formation of the Contract.
	31.3	Upon the successful Bidder's furnishing of the performance security pursuant to ITB Clause 33, the Procuring agency will promptly notify each unsuccessful Bidder and will discharge its bid security, pursuant to ITB Clause 15.
<b>32. Signing of Contract</b>	32.1	At the same time as the Procuring agency notifies the successful Bidder that its bid has been accepted, the Procuring agency will send the Bidder the Contract Form provided in the bidding documents, incorporating all agreements between the parties.

	32.2	Within thirty (30) days of receipt of the Contract Form, the successful Bidder shall sign and date the contract and return it to the Procuring
--	------	--

		agency.
<b>33 Performance Security</b>	33.1	Within twenty (20) days of the receipt of notification of award from the Procuring agency, the successful Bidder shall furnish the performance security in accordance with the Conditions of Contract, in the Performance Security Form provided in the bidding documents, or in another form acceptable to the Procuring agency.
	33.2	Failure of the successful Bidder to comply with the requirement of ITB Clause 32 or ITB Clause 33.1 shall constitute sufficient grounds for the annulment of the award and forfeiture of the bid security, in which event the Procuring agency may make the award to the next lowest evaluated Bidder or call for new bids.
<b>34. Corrupt or Fraudulent Practices</b>	34.1	<p>The Government of Khyber Pakhtunkhwa requires that Procuring agency's (including beneficiaries of donor agencies' loans), as well as Bidders/Suppliers/Contractors under Government-financed contracts, observe the highest standard of ethics during the procurement and execution of such contracts. In pursuance of this policy, the KPPRA, in accordance with the KPP Act, 2009 and Rules made thereunder:</p> <ul style="list-style-type: none"> <li>a. defines, for the purposes of this provision, the terms set forth below as follows: <ul style="list-style-type: none"> <li>i. "corrupt practice" means the offering, giving, receiving or soliciting of anything of value to influence the action of a public official in the procurement process or in contract execution; and</li> <li>ii. "fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Procuring agency, and includes collusive practice among Bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to deprive the Procuring agency of the benefits of free and open competition;</li> </ul> </li> <li>b. will reject a proposal for award if it determines that the Bidder recommended for award has engaged in corrupt or fraudulent practices in competing for the contract in question;</li> <li>c. will declare a firm ineligible, either indefinitely or for a stated period of time, to be awarded a Government-financed contract if it at any time determines that the firm has engaged in corrupt or fraudulent practices in competing for, or in executing, a Government-financed contract.</li> </ul>
	34.2	Furthermore, Bidders shall be aware of the provision stated in sub-clause 5.4 and sub-clause 24.1 of the General Conditions of Contract.
<b>36. Integrity Pact</b>	35.1	The Bidder shall sign and stamp the Integrity Pact provided at Form - 7 to Bid in the Bidding Document for all Provincial Government procurement contracts exceeding Rupees ten million. Failure to such Integrity Pact shall make the bidder non-responsive.

## **Part One - Section II.**

### **General Conditions of Contract**

## **Notes on the General Conditions of Contract (GCC)**

The General Conditions of Contract in Part One Section II, read in conjunction with the Special Conditions of Contract in Part Two Section III and other documents listed therein, should be a complete document expressing all the rights and obligations of the parties.

The General Conditions of Contract herein shall not be altered. Any changes and complementary information, which may be needed, shall be introduced only through the Special Conditions of Contract in Part Two Section III.

## Table of Clauses

1.	Definitions	26
2.	Application	26
3.	Country of Origin	26
4.	Standards	27
5.	Use of Contract Documents and Information; Inspection and Audit by the Bank	27
6.	Patent Rights	27
7.	Performance Security	27
8.	Inspections and Tests	28
9.	Packing	28
10.	Delivery and Documents	29
11.	Insurance	29
12.	Transportation	29
13.	Incidental Services	29
14.	Spare Parts	29
15.	Warranty	30
16.	Payment	30
17.	Prices	30
18.	Change Orders	31
19.	Contract Amendments	31
20.	Assignment	31
21.	Subcontracts	31
22.	Delays in the Supplier's Performance	31
23.	Liquidated Damages	31
24.	Termination for Default	32
25.	Force Majeure	32
26.	Termination for Insolvency	33
27.	Termination for Convenience	33
28.	Resolution of Disputes	33
29.	Governing Language	33
30.	Applicable Law	34
31.	Notices	34
32.	Taxes and Duties	34

## General Conditions of Contract

<b>1. Definitions</b>	1.1	<p>In this Contract, the following terms shall be interpreted as indicated:</p> <ul style="list-style-type: none"> <li>a. “The Contract” means the agreement entered into between the Procuring agency and the Supplier, as recorded in the Contract Form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.</li> <li>b. “The Contract Price” means the price payable to the Supplier under the Contract for the full and proper performance of its contractual obligations.</li> <li>c. “The Goods” means all of the equipment, machinery, and/or other materials which the Supplier is required to supply to the Procuring agency under the Contract.</li> <li>d. “The Services” means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training, and other such obligations of the Supplier covered under the Contract.</li> <li>e. “GCC” means the General Conditions of Contract contained in this section.</li> <li>f. “SCC” means the Special Conditions of Contract.</li> <li>g. “The Procuring agency” means the organization purchasing the Goods, as named in SCC.</li> <li>h. “The Procuring agency’s country” is the country named in SCC.</li> <li>i. “The Supplier” means the individual or firm supplying the Goods and Services under this Contract.</li> <li>j. “The Project Site,” where applicable, means the place or places named in SCC.</li> <li>k. “Day” means calendar day.</li> </ul>
<b>2. Application</b>	2.1	These General Conditions shall apply to the extent that they are not superseded by provisions of other parts of the Contract.
<b>3. Country of Origin</b>	3.1	All Goods and Services supplied under the Contract shall have their origin in the countries and territories eligible under the rules and further elaborated in the SCC.
	3.2	For purposes of this Clause, “origin” means the place where the Goods were mined, grown, or produced, or from which the Services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized new product results that is substantially different in basic characteristics or in purpose or utility from its components.
	3.3	The origin of Goods and Services is distinct from the nationality of the Supplier.

<b>4. Standards</b>	4.1	The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications, and, when no applicable standard is mentioned, to the authoritative standards appropriate to the Goods' country of origin. Such standards shall be the latest issued by the concerned institution.
<b>5. Use of Contract Documents and Information; Inspection and Audit by the Government</b>	5.1	The Supplier shall not, without the Procuring agency's prior written consent, disclose the Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the Procuring agency in connection therewith, to any person other than a person employed by the Supplier in the performance of the Contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.
	5.2	The Supplier shall not, without the Procuring agency's prior written consent, make use of any document or information enumerated in GCC Clause 5.1 except for purposes of performing the Contract.
	5.3	Any document, other than the Contract itself, enumerated in GCC Clause 5.1 shall remain the property of the Procuring agency and shall be returned (all copies) to the Procuring agency on completion of the Supplier's performance under the Contract if so required by the Procuring agency.
	5.4	The Supplier shall permit the Procuring agency 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 procuring agency, if so required.
<b>6. Patent Rights</b>	6.1	The Supplier shall indemnify the Procuring agency against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the Goods or any part thereof in the Procuring agency's country.
<b>7. Performance Security</b>	7.1	Within twenty (20) days of receipt of the notification of Contract award, the successful Bidder shall furnish to the Procuring agency the performance security in the amount specified in SCC.
	7.2	The proceeds of the performance security shall be payable to the Procuring agency as compensation for any loss resulting from the Supplier's failure to complete its obligations under the Contract.
	7.3	The performance security shall be denominated in the currency of the Contract acceptable to the Procuring agency and shall be in one of the following forms:  a. a bank guarantee or an irrevocable letter of credit issued by a reputable bank located in the Procuring agency's country, in the form provided in the bidding documents or another form acceptable to the Procuring agency; or  b. a cashier's or certified check.

	7.4	The performance security will be discharged by the Procuring agency 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 SCC.
<b>8. Inspections and Tests</b>	8.1	The Procuring agency or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Contract specifications at no extra cost to the Procuring agency. SCC and the Technical Specifications shall specify what inspections and tests the Procuring agency requires and where they are to be conducted. The Procuring agency shall notify the Supplier in writing, in a timely manner, of the identity of any representatives retained for these purposes.
	8.2	The inspections and tests may be conducted on the premises of the Supplier or its subcontractor(s), at point of delivery, and/or at the Goods' final destination. If conducted on the premises of the Supplier or its subcontractor(s), all reasonable facilities and assistance, including access to drawings and production data, shall be furnished to the inspectors at no charge to the Procuring agency.
	8.3	Should any inspected or tested Goods fail to conform to the Specifications, the Procuring agency may reject the Goods, and the Supplier shall either replace the rejected Goods or make alterations necessary to meet specification requirements free of cost to the Procuring agency.
	8.4	The Procuring agency's right to inspect, test and, where necessary, reject the Goods after the Goods' arrival in the Procuring agency's country shall in no way be limited or waived by reason of the Goods having previously been inspected, tested, and passed by the Procuring agency or its representative prior to the Goods' shipment from the country of origin.
	8.5	Nothing in GCC Clause 8 shall in any way release the Supplier from any warranty or other obligations under this Contract.
<b>9. Packing</b>	9.1	The Supplier shall provide such packing of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the Contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit.
	9.2	The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified in SCC, and in any subsequent
		Instructions ordered by the Procuring agency.

<b>10. Delivery and Documents</b>	10.1	Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in the Schedule of Requirements. The details of shipping and/or other documents to be furnished by the Supplier are specified in SCC.
	10.2	Documents to be submitted by the Supplier are specified in SCC.
<b>11. Insurance</b>	11.1	The Goods supplied under the Contract shall be delivered duty paid (DDP) under which risk is transferred to the buyer after having been delivered; hence insurance coverage is seller's responsibility.
<b>12. Transportation</b>	12.1	The Supplier is required under the Contract to transport the Goods to a specified place of destination within the Procuring agency's country, transport to such place of destination in the Procuring agency's country, including insurance and storage, as shall be specified in the Contract, shall be arranged by the Supplier, and related costs shall be included in the Contract Price.
<b>13. Incidental Services</b>	13.1	<p>The Supplier may be required to provide any or all of the following services, including additional services, if any, specified in SCC:</p> <ul style="list-style-type: none"> <li>a. performance or supervision of on-site assembly and/or start-up of the supplied Goods;</li> <li>b. furnishing of tools required for assembly and / or maintenance of the supplied Goods;</li> <li>c. furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied Goods;</li> <li>d. performance or supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties, provided that this service shall not relieve the Supplier of any warranty obligations under this Contract; and</li> <li>e. training of the Procuring agency's personnel, at the Supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied Goods.</li> </ul>
	13.2	Prices charged by the Supplier for incidental services, if not included in the Contract Price for the Goods, shall be agreed upon in advance by the parties and shall not exceed the prevailing rates charged for other parties by the Supplier for similar services.
<b>14. Spare Parts</b>	14.1	<p>As specified in SCC, the Supplier may be required to provide any or all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier:</p> <ul style="list-style-type: none"> <li>a. such spare parts as the Procuring agency may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under the Contract; and</li> <li>b. in the event of termination of production of the spare parts: <ul style="list-style-type: none"> <li>i. advance notification to the Procuring agency of the pending termination, in sufficient time to permit the Procuring agency to procure needed requirements;</li> <li>ii. Following such termination, furnishing at no cost to the Procuring agency, the blueprints, drawings, and specifications of the spare parts, if requested.</li> </ul> </li> </ul>

<b>15. Warranty</b>	15.1	The Supplier warrants that the Goods supplied under the Contract are new, unused, of the most recent or current models, and that they incorporate all recent improvements in design and materials unless provided otherwise in the Contract. The Supplier further warrants that all Goods supplied under this Contract shall have no defect, arising from design, materials, or workmanship (except when the design and/or material is required by the Procuring agency's specifications) or from any act or omission of the Supplier, that may develop under normal use of the supplied Goods in the conditions prevailing in the country of final destination.
	15.2	This warranty shall remain valid for twelve (12) months after the Goods, or any portion thereof as the case may be, have been delivered to and accepted at the final destination indicated in the Contract, or for eighteen (18) months after the date of shipment from the port or place of loading in the source country, whichever period concludes earlier, unless specified otherwise in SCC.
	15.3	The Procuring agency shall promptly notify the Supplier in writing of any claims arising under this warranty.
	15.4	Upon receipt of such notice, the Supplier shall, within the period specified in SCC and with all reasonable speed, repair or replace the defective Goods or parts thereof, without costs to the Procuring agency.
	15.5	If the Supplier, having been notified, fails to remedy the defect(s) within the period specified in SCC, within a reasonable period, the Procuring agency may proceed to take such remedial action as may be necessary, at the Supplier's risk and expense and without prejudice to any other rights which the Procuring agency may have against the Supplier under the Contract.
<b>16. Payment</b>	16.1	The method and conditions of payment to be made to the Supplier under this Contract shall be specified in SCC.
	16.2	The Supplier's request(s) for payment shall be made to the Procuring agency in writing, accompanied by an invoice describing, as appropriate, the Goods delivered and Services performed, and by documents submitted pursuant to GCC Clause 10, and upon fulfillment of other obligations stipulated in the Contract.
	16.3	Payments shall be made promptly by the Procuring agency, but in no case later than sixty (60) days after submission of an invoice or claim by the Supplier.
	16.4	The currency of payment is Pak. Rupees.
<b>17. Prices</b>	17.1	Prices charged by the Supplier for Goods delivered and Services performed under the Contract shall not vary from the prices quoted by the Supplier in its bid, with the exception of any price adjustments authorized in SCC or in the Procuring agency's request for bid validity extension, as the case may be.

<b>18. Change Orders</b>	18.1	<p>The Procuring agency may at any time, by a written order given to the Supplier pursuant to GCC Clause 31, make changes within the general scope of the Contract in any one or more of the following:</p> <ul style="list-style-type: none"> <li>a. drawings, designs, or specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the Procuring agency;</li> <li>b. the method of shipment or packing;</li> <li>c. the place of delivery; and/or</li> <li>d. the Services to be provided by the Supplier.</li> </ul>
	18.2	<p>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 delivery 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 Procuring agency's change order.</p>
<b>19. Contract Amendments</b>	19.1	<p>Subject to GCC Clause 18, no variation in or modification of the terms of the Contract shall be made except by written amendment signed by the parties.</p>
<b>20. Assignment</b>	20.1	<p>The Supplier shall not assign, in whole or in part, its obligations to perform under this Contract, except with the Procuring agency's prior written consent.</p>
<b>21. Subcontracts</b>	21.1	<p>The Supplier shall notify the Procuring agency in writing of all subcontracts awarded under this Contract if not already specified in the bid. Such notification, in the original bid or later, shall not relieve the Supplier from any liability or obligation under the Contract.</p>
	21.2	<p>Subcontracts must comply with the provisions of GCC Clause 3.</p>
<b>22. Delays in the Supplier's Performance</b>	22.1	<p>Delivery of the Goods and performance of Services shall be made by the Supplier in accordance with the time schedule prescribed by the Procuring agency in the Schedule of Requirements.</p>
	22.2	<p>If at any time during performance of the Contract, the Supplier or its subcontractor(s) should encounter conditions impeding timely delivery of the Goods and performance of Services, the Supplier shall promptly notify the Procuring agency in writing of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the Supplier's notice, the Procuring agency shall evaluate the situation and may at its discretion extend the Supplier's time for performance, with or without liquidated damages, in which case the extension shall be ratified by the parties by amendment of Contract.</p>
	22.3	<p>Except as provided under GCC Clause 25, a delay by the Supplier in the performance of its delivery obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC Clause 23, unless an extension of time is agreed upon pursuant to GCC Clause 22.2 without the application of liquidated damages.</p>
<b>23. Liquidated Damages</b>	2.31	<p>Subject to GCC Clause 25, if the Supplier fails to deliver any or all of the Goods or to perform the Services within the period(s) specified in</p>

		the Contract, the Procuring agency shall, without prejudice to its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to the percentage specified in SCC of the delivered price of the delayed Goods or unperformed Services for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the percentage specified in SCC. Once the maximum is reached, the Procuring agency may consider termination of the Contract pursuant to GCC Clause 24.
<b>24. Termination for Default</b>	24.1	<p>The Procuring agency, without prejudice to any other remedy for breach of Contract, by written notice of default sent to the Supplier, may terminate this Contract in whole or in part:</p> <ul style="list-style-type: none"> <li>a. if the Supplier fails to deliver any or all of the Goods within the period(s) specified in the Contract, or within any extension thereof granted by the Procuring agency pursuant to GCC Clause 22; or</li> <li>b. if the Supplier fails to perform any other obligation(s) under the Contract.</li> <li>c. if the Supplier, in the judgment of the Procuring agency has engaged in corrupt or fraudulent practices in competing for or in executing the Contract.</li> </ul> <p>For the purpose of this clause:</p> <p>“corrupt practice” means the offering, giving, receiving or soliciting of anything of value to influence the action of a public official in the procurement process or in contract execution.</p> <p>“fraudulent practice” means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Borrower, and includes collusive practice among Bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to deprive the Borrower of the benefits of free and open competition.</p>
	24.2	In the event the Procuring agency terminates the Contract in whole or in part, pursuant to GCC Clause 24.1, the Procuring agency may procure, upon such terms and in such manner as it deems appropriate, Goods or Services similar to those undelivered, and the Supplier shall be liable to the Procuring agency for any excess costs for such similar Goods or Services. However, the Supplier shall continue performance of the Contract to the extent not terminated.
<b>25. Force Majeure</b>	25.1	Notwithstanding the provisions of GCC Clauses 22, 23, and 24, 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.

	25.2	For purposes of this clause, “Force Majeure” means an event beyond the control of the Supplier and not involving the Supplier’s fault or negligence and not foreseeable. Such events may include, but are not restricted to, acts of the Procuring agency in its sovereign capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions, and freight embargoes.
	25.3	If a Force Majeure situation arises, the Supplier shall promptly notify the Procuring agency in writing of such condition and the cause thereof. Unless otherwise directed by the Procuring agency 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.
<b>26. Termination for Insolvency</b>	26.1	The Procuring agency may at any time terminate the Contract by giving written notice to the Supplier if the Supplier becomes bankrupt or otherwise insolvent. In this event, termination will be without compensation to the Supplier, provided that such termination will not prejudice or affect any right of action or remedy which has accrued or will accrue thereafter to the Procuring agency.
<b>27. Termination for Convenience</b>	27.1	The Procuring agency, 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 Procuring agency’s convenience, the extent to which performance of the Supplier under the Contract is terminated, and the date upon which such termination becomes effective.
	27.2	The Goods that are complete and ready for shipment within thirty (30) days after the Supplier’s receipt of notice of termination shall be accepted by the Procuring agency at the Contract terms and prices. For the remaining Goods, the Procuring agency may elect: <ul style="list-style-type: none"> <li>a. to have any portion completed and delivered at the Contract terms and prices; and/or</li> <li>b. to cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Services and for materials and parts previously procured by the Supplier.</li> </ul>
<b>28. Resolution of Disputes</b>	28.1	The Procuring agency 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.
	28.2	If, after thirty (30) days from the commencement of such informal negotiations, the Procuring agency and the Supplier have been unable to resolve amicably a Contract dispute, either party may require that the dispute be referred for resolution to the formal mechanisms specified in SCC. These mechanisms may include, but are not restricted to, conciliation mediated by a third party, adjudication in an agreed manner and/or arbitration.
<b>29. Governing Language</b>	29.1	The Contract shall be written in the language specified in SCC. Subject to GCC Clause 30, the version of the Contract written in the specified language shall govern its interpretation. All correspondence and other documents pertaining to the Contract which are exchanged by the parties shall be written in the same language.

<b>30. Applicable Law</b>	30.1	The Contract shall be interpreted in accordance with the laws of the Procuring agency's country, unless otherwise specified in SCC.
<b>31. Notices</b>	31.1	Any notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing or by cable, telex, or facsimile and confirmed in writing to the other party's address specified in SCC.
	31.2	A notice shall be effective when delivered or on the notice's effective date, whichever is later.
<b>32. Taxes and Duties</b>	32.1	Supplier shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted Goods to the Procuring agency.



**Government of Khyber Pakhtunkhwa**

**Health Department**

**Directorate General Health Services  
Khyber Pakhtunkhwa Peshawar**

## **Standard Bidding Documents**

**For National Competitive Bidding  
Pakistan**

**For**

**SELECTION AND RATE CONTRACTING OF DRUGS / MEDICINES,  
MEDICAL DEVICES, SURGICAL DISPOSABLES &  
NON-DRUG ITEMS**

**FOR THE FINANCIAL YEARS 2021-23**

**MEDICINE COORDINATION CELL (MCC)**

**APRIL 2021**

## **PART TWO (PROCUREMENT SPECIFIC PROVISIONS)**

- Invitation for Bids (IFB)
- Bid Data Sheet (BDS)
- Special Conditions of Contract (SCC)
- Schedule of Requirements
- Technical Specifications
- Sample Forms
- Eligibility

## Preface

These Bidding Documents have been prepared for use by procuring agencies in the procurement of goods through National Competitive Bidding (NCB).

In order to simplify the preparation of bidding documents for each procurement, the Bidding Documents are grouped in two parts based on provisions which are fixed and that which are specific for each procurement. Provisions which are intended to be used unchanged are in Part one, which includes Section I, Instructions to Bidders, and Section II, General Conditions of Contract. Data and provisions specific to each procurement and contract are included in Part Two which includes Section II, Bid Data Sheet; Section III, Special Conditions of Contract; Section IV, Schedule of Requirements; Section V, Technical Specifications; and the forms to be used in Section I, Invitation for Bids, and Section VI, Sample Forms.

This is Part Two and contains data and provisions specific to each procurement. Care should be taken to check the relevance of the provisions of the Bidding Documents against the requirements of the specific goods to be procured. The following general directions should be observed when using the documents. In addition, each section is prepared with notes intended only as information for the Procuring agency or the person drafting the bidding documents. They shall not be included in the final documents, except for the notes introducing Section VI, Forms, where the information is useful for the Bidder.

- a. Specific details, such as the “name of the Procuring agency” and “address for bid submission,” should be furnished in the Invitation for Bids, in the Bid Data Sheet, and in the Special Conditions of Contract. The final documents should contain neither blank spaces nor options.
- b. Amendments, if any, to the Instructions to Bidders and to the General Conditions of Contract should be made through the Bid Data Sheet and the Special Conditions of Contract, respectively.
- c. Footnotes or notes in italics included in the Invitation for Bids, Bid Data Sheet, Special Conditions of Contract, and in the Schedule of Requirements are not part of the text of the document, although they contain instructions that the Procuring agency should strictly follow. The final document should contain no footnotes.
- d. The criteria for bid evaluation and the various methods of evaluation in the Instructions to Bidders (Clauses 25.3 and 25.4, respectively) should be carefully reviewed. Only those that are selected to be used for the procurement in question should be retained and expanded, as required, in the Bid Data Sheet or in the Technical Specifications, as appropriate. The criteria that are not applicable should be deleted from the Bid Data Sheet.
- e. Clauses included in the Special Conditions of Contract are illustrative of the provisions that should be drafted specifically by the Procuring agency for each procurement.
- f. The forms provided in Section VI should be completed by the Bidder or the Supplier; the footnotes in these forms should remain, since they contain instructions which the Bidder or the Supplier should follow.

## Table of Contents - Part Two

<b>Contents</b>	<b>Page No.</b>
Section I. Invitation for Bids	39
Section II. Bid Data Sheet	42
Section III. Special Conditions of Contract	47
Table of clauses	48
Section IV. Schedule of Requirements (SOR)	54
Section V. Technical Specifications	146
Section VI. Sample Forms	157
Sample Forms	166
1. Bid Cover Sheet Bid Form-1	166
2. Letter of Intention Bid Form- 2	169
3. Affidavit Bid Form-3	170
4. Price Schedule Format Bid Form -4	171
5. Integrity Pact Bid Form-5	172
6. MCC Rate Contract Agreement Bid Form-6	173

## Part Two

### Section I. Invitation for Bids

#### Notes on the Invitation for Bids

The Invitation for Bids (IFB) has been issued as an advertisement in leading newspapers of general circulation in the Province of Khyber Pakhtunkhwa as well as on the web site of the Health Department ([www.healthkp.gov.pk](http://www.healthkp.gov.pk)) and ([www.dghskp.gov.pk](http://www.dghskp.gov.pk)) by allowing at least fifteen days for NCB for bid preparation and submission.

The Invitation for Bids provides information that enables interested bidders to decide whether to participate. Apart from the essential items listed in the Standard Bidding Documents (SBD), the Invitation for Bids also indicates the important bid evaluation criteria or qualification requirement (for example, a requirement for a minimum level of experience in manufacturing a similar type of goods for which the Invitation for Bids is issued) so that the bidders should give their best and final prices as no negotiations are allowed.

The Invitation for Bids is incorporated into these Standard Bidding Documents (SBDs). The information contained in the Invitation for Bids (IFB) conforms to the bidding documents and in particular to the relevant information in the Bid Data Sheet.

## Invitation for Bids

**Government Medicine Coordination Cell  
Directorate General Health Services  
Khyber Pakhtunkhwa, Peshawar**

**SELECTION AND RATE CONTRACTING (CONTRACT FRAME WORK  
AGREEMENT) OF DRUGS / MEDICINES, MEDICAL DEVICES, SURGICAL  
DISPOSABLES & NON-DRUG ITEMS  
FOR THE YEARS 2021-23**

1. In compliance with the Khyber Pakhtunkhwa Public Procurement Act-2012 and Khyber Pakhtunkhwa Procurement Regulatory Authority (KPPRA) Rules-2014, Government Medicine Coordination Cell (Government MCC), Directorate General Health Services Khyber Pakhtunkhwa, Warsak Road, Peshawar invites sealed bids from i) Manufacturer/s and / or Importer/s of drugs / medicines authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan, registered as such with the Drug Regulatory Authority of Pakistan (DRAP) for the quoted item/s falling under The Drug Act 1976 & Rules framed there under; and ii) Manufacturer/s of Medical Devices in Pakistan, registered as such with the DRAP for the quoted item/s and regulated under the DRAP Act 2012 and the Rules framed thereunder; and iii) Importer/s of Medical Devices, duly authorized by the goods' Principal Manufacturer or producer to import / supply the said goods in Pakistan, as registered and regulated as such for the quoted item/s under the DRAP Act 2012 and Rules framed thereunder; and iv) Manufacturer/s of Non-Drug Items (NDIs) in Pakistan; and v) Importer/s of NDIs, duly authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan.
2. Manufacturer/s and/or Importer/s of various items interested to enter in this bidding competition must obtain separate application form from the office of the Director General Drug Control and Pharmacy Services (Govt. MCC), Old FATA Secretariat Warsak Road Peshawar during office hours on any working day till **26<sup>th</sup> April 2021**. At the time of submission of the bid, the original receipt of non-refundable cash payment of Pak Rupees Five Thousand (Rs. 5000/-) per application form shall be submitted with technical bid. No Application Form shall be issued after 2:00 Pm, **26<sup>th</sup> April 2021**.
3. Bidding competition under this advertisement shall be conducted through **Single Stage-Two Envelopes Bidding Procedure** as per KPPRA Act 2012 and Rules frame there under. Under this procedure, the bidders should submit the bids in two sealed envelopes of technical and financial bids, each of which must bear on them the clearly written words 'Government MCC Technical Bid 2021-23' and 'Government MCC Financial Bid 2021-23' as well as the full and complete identification of the bidder along with its postal and email addresses and phone number/s on each of the respective envelope. Both these sealed and labeled envelopes should be placed inside another outer envelope of appropriate size which should also be sealed and should bear the clearly written words "Bid for Govt. MCC 2021-23 along with the identification and contact details of the bidder.
4. The Standard Bidding Documents, other than the application form mentioned above, for this bidding competition may be downloaded from [www.healthkp.gov.pk](http://www.healthkp.gov.pk) and [www.dghskp.gov.pk](http://www.dghskp.gov.pk)
5. A Pre bid meeting is scheduled to be held on 12<sup>th</sup> and 13<sup>th</sup> April 2021, at 11:00 am, at the Committee room of Directorate General Health Services Khyber Pakhtunkhwa Old FATA Secretariat Warsak Road Peshawar in the following groups.
  - a. 12<sup>th</sup> April 2021- For bidders (Manufacturers and Importers of General Medicines, IV Fluids, Biologicals and Dry Powder Injectables).

- b. 13<sup>th</sup> April 2021- For bidders (Manufacturers and Importers of Medical Devices, Non Drug Items, Cardiac Stents, Cotton and Related Goods)

The bidders shall thoroughly study the standard bidding documents (SBDs) before the Pre-Bid meeting and bring their query (s)/suggestion(s) to the forum for clarification/understanding and shall be submitted in written on or before the Pre-Bid Meeting. In case of non-submission as hard copy on or before the meeting day, the query(ies)/suggestion(s) shall not be considered/entertained on the day of meeting or afterwards.

6. Bidders must submit sealed bids to the office of Director General Drug Control and Pharmacy Services, Old FATA Secretariat Warsak Road Peshawar **on or before 11:00 a.m. sharp on 27<sup>th</sup> April 2021**. Any bids presented / submitted / received later than this deadline, or delivered to some office other than the above office, shall not be considered and shall be rejected without any further processing.
7. Mandatory Bid Security / Earnest Money amounting to a flat rate of Rupees Eight Hundred Thousands only (Rs.800, 000/-) from each bidder in the shape of **Call Deposit Receipt (CDR)** in the name of the Director General Health Services, Khyber Pakhtunkhwa is required to be submitted in original along with the Financial Bid within its sealed envelope and shall be from the account of the firm/manufacturer/importer. A separate photocopy of this Bids Security financial instrument should also be placed inside the sealed envelope of Technical Proposal. Ordinary crossed or open Cheques shall not be acceptable as Bid's security.
8. Quotation must be computer typed & printed; the Offered rate, Trade Price (TP) and Maximum Retail Price (MRP) must be written both in words & figures. All pages of the submitted bid shall be signed, numbered and duly stamped by the authorized person of the bidding entity as mentioned in the SBDs.
9. The bidders are required to submit the unit prices (Offered, TP and MRP) of quoted items on the format as prescribed for financial bid in the Standard Bidding Documents.
10. Quotations with cutting, erasing, and over-writing shall not be accepted to the extent of that particular quoted item.
11. To facilitate the data entry during bids processing, all bidders are required to submit the quoted product list as per the prescribed proformas in the approved Standard Bidding Documents for this bidding competition, in soft form in MS Excel format (and not in other software formats or images) on computer CD/DVD, duly labeled by a permanent marker with the name of bidder firm along with the words 'Government MCC 2021-23. The bidders must ensure that said computer CD/DVD or USB is open able and readable. Moreover, in the same context, the bidders are also required to submit a hard copy of Tape bind booklet bid, having table of contents (Indexing with proper page number and contents mentioned in the start of bid and each page of the submitted bid shall be properly numbered, signed and stamped by the authorized person of the bidding entity).
12. Bidders are required and encouraged to offer the most competitive lowest price/s of their quoted item/s as no negotiations on quoted price/s shall be allowed under the rules.
13. Bids will In sha Allah be opened by the Technical & Evaluation Committee of Government MCC at **11:30 a.m. (Sharp) on 27<sup>th</sup> APRIL 2021** in the office of Directorate General Drug Control & Pharmacy Services, Warsak Road, Old FATA Secretariat Peshawar in the presence of bidders or their representatives (who choose to attend the bids opening process).
14. The Directorate General Health Services reserves the right to reject any or all the bids under clause 47 of KPPRA Procurement Rules 2014.

Director General Health Services Khyber Pakhtunkhwa,  
Old FATA Secretariat Warsak Road, Peshawar  
**Tel No: 091-9222824, 091-9211702,**  
**Email: mccdgcps@gmail.com**

## Section II. Bid Data Sheet

### BID DATA SHEET

ITB Ref.	Introduction/Description	Detail
ITB 1.1	Name of Procuring Agency of Government of Khyber Pakhtunkhwa.	Director General Health Services, Khyber Pakhtunkhwa as the overall head of the procuring entity, Health Department Government of Khyber Pakhtunkhwa.
ITB 1.1	Loan or credit or Project allocation number. Loan or credit or Project allocation amount.	Not Applicable
ITB 1.1	Name of Project	Not Applicable
ITB 1.1	Name of Contract	Not Applicable
ITB 4.1	Name of Procuring agency.	Director General Health Services, Khyber Pakhtunkhwa as the overall head of the procuring entity, Health Department Government of Khyber Pakhtunkhwa.
ITB 6.1	Procuring agency's address, telephone, telex, and facsimile, numbers.	Director General Health Services Khyber Pakhtunkhwa Peshawar Tel No: 091- 9214084, 091-9210269 Fax No: 091- 9210230 Email: <a href="mailto:mccdgdcp@gmail.com">mccdgdcp@gmail.com</a>
ITB 8.1	Language of the bid.	English
<b>Bid Price and Currency</b>		
ITB 11.2	Price quoted shall be:	Pakistani Rupees (Rs.)
ITB 11.5	The price shall be fixed	The price shall be fixed and valid till 30 <sup>th</sup> June 2023.
<b>Preparation and Submission of Bids</b>		
ITB 13.3 (d)	Qualification requirements.	<p><b>I.</b> Manufacturer/s and / or Importer/s of drugs / medicines authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan, registered as such with the Drug Regulatory Authority of Pakistan (DRAP) for the quoted item/s falling under The Drug Act 1976 &amp; Rules framed there under; and</p> <p><b>II.</b> <b>Manufacturer</b> of Medical Devices in Pakistan, registered as such with the DRAP for the quoted item/s and regulated under the DRAP Act 2012 and the Rules framed there under; and</p> <p><b>III.</b> <b>Importer</b> of Medical Devices, duly authorized by the goods' Principal Manufacturer or producer to import / supply the said goods in Pakistan, as registered and regulated as such for the</p>

		quoted item/s under the DRAP Act 2012 and Rules framed there under; and <b>V. Manufacturer</b> of Non-Drug Items (NDIs) in Pakistan; and <b>V. Importer</b> of NDIs, duly authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan.
<b>ITB 14.3 (b)</b>	Spare parts required for ----- of years of operation	Not Applicable
<b>ITB 15.1</b>	Amount of bid security.	Rs. 800,000/-
<b>ITB 16.1</b>	Bid validity period.	180 days from the date of opening of bids
<b>ITB 17.1</b>	Number of copies.	One (ORIGINAL BID)
<b>ITB 18.2 (a)</b>	Address for bid submission.	Directorate General Drug Control and Pharmacy services, Old FATA Secretariat, Warsak Road, Peshawar
<b>ITB 18.2 (b)</b>	IFB title and number.	Selection and Rate Contracting (Contract Framework Agreement) of Drugs / Medicines, Medical Devices, Surgical Disposables & Non-Drug items for the year 2021-23.
<b>ITB 19.1</b>	Deadline for bid submission.	<b>Before or up to 11:00 a.m. sharp on 27<sup>th</sup> April 2021.</b>
<b>ITB 22.1</b>	Time, Date and Place for bid opening.	<b>11:30 hours on 27<sup>th</sup> April 2021 in the Conference Room of Directorate General Drug Control and Pharmacy services, Old FATA Secretariat, Warsak Road, Peshawar</b>
<b>Bid Evaluation</b>		
<b>ITB 25.3</b>	Criteria for bid evaluation.	Merit Point Evaluation (Highest Ranking Fair Bid). The items ranked highest in merit points (obtained through, and based on, technical and financial evaluation) will get unit rate central contract. (Section-V of these SBDs).
<b>ITB 25.4 (a)</b>	One option only	Not Applicable
<b>ITB 25.4 (b)</b>	Delivery schedule. Relevant parameters in accordance with option selected.	
<b>Option I</b> <b>Option II</b> <b>Option III</b>	Adjustment expressed as a percentage, or adjustment expressed in an amount in the currency of bid evaluation, or adjustment expressed in an amount in the currency of bid evaluation.	Not Applicable
<b>ITB 25.4 (c)(ii)</b>	Deviation in payment schedule. Annual interest rate.	Not Applicable
<b>ITB 25.4 (d)</b>	Cost of spare parts.	Not Applicable
<b>ITB 25.4 (e)</b>	Spare parts and after sales service facilities in the Procuring agency's country.	Not Applicable
<b>ITB 25.4 (f)</b>	Operating and maintenance costs.	Not Applicable
<b>ITB 25.4 (g)</b>	Performance and productivity of equipment.	Not Applicable

<b>ITB 25.4 (h)</b>	Details on the evaluation method or reference to the Technical Specifications	As in section on Technical Evaluation of bids. The evaluation parameters of the
---------------------	---	---

		<p>quoted item/s may include, but not limited to, any or all of the methods including physical inspection, examination, testing/using by the end user/s and or laboratory testing and/ or market survey against any parameter/s, as deemed appropriate by the procuring Agency or any of its committees or sub-committees. Any discrepancy found during the market survey shall lead to disqualification of the firm/product (s).</p> <p>Physical Inspection of manufacturers and importers will be carried out through a uniform checklist/Performa.</p> <p>All the certifications from accredited bodies, as the case may be, shall contain the quoted product (s) in its scope, moreover the accredited body shall be authorized to certify the quoted product (s).</p> <p>In case of products having Multiple APIs/Raw material the marks for GD, CoA, APIs or Raw material Source accreditation will be awarded only where these documents are submitted for all ingredients/components of the quoted products For Example. Sitagliptin + Metformin, IV Cannula (Plastic and Needle etc.)</p> <p>In case the Supplier had been awarded marks in product evaluation parameter during the technical evaluation for API source accreditation for Drugs / Medicines, and for medical grade material certification for medical devices &amp; Non-Drug Items, and for Pharmaceutical grade certification for immediate containers of Drugs/medicines shall warranty the supply of all such goods with the same certified quality, material and specification/s to the Purchasing Agency/ies throughout the validity period of contract agreement.</p>
<b>ITB 25.4 alternative</b>	Specify the evaluation factors.	Not Applicable
<b>Contract Award</b>		

<b>ITB 29.1</b>	Percentage for quantity increase or decrease.	The Procuring Agency in the capacity of being the overall head of the Government Medicine Coordination Cell, or otherwise has the authority to regulate, if deemed appropriate, under the provisions in ITB 29.1 through imposing restrictions and / or classifying and / or grouping any selected quoted item/s for stopping, increasing or decreasing the purchase of such item/s by the Purchasing Agency/ies to rationalize and / or control the use and / or misuse of such item/s.
-----------------	---	--

## **Section III. Special Conditions of Contract**

### **Notes on the Special Conditions of Contract**

Similar to the Bid Data Sheet in Section II, the clauses in this Section are intended to assist the Procuring agency in providing contract-specific information in relation to corresponding clauses in the General Conditions of Contract.

The provisions of Section III complement the General Conditions of Contract included in Part one, Section II, specifying contractual requirements linked to the special circumstances of the Procuring agency, the Procuring agency's country, the sector, and the Goods purchased. In preparing Section III, the following aspects should be checked:

- a. Information that complements provisions of Part one Section II must be incorporated.
- b. Amendments and/or supplements to provisions of Part one Section II, as necessitated by the circumstances of the specific purchase, must also be incorporated.

### Section III. Special Conditions of Contract

**Table of Clauses**

1.	DEFINITIONS (GCC CLAUSE 1)	49
2.	COUNTRY OF ORIGIN (GCC CLAUSE 3)	50
3.	PERFORMANCE SECURITY (GCC CLAUSE 7)	50
4.	INSPECTIONS AND TESTS (GCC CLAUSE 8)	50
5.	PACKING (GCC CLAUSE 9)	51
6.	DELIVERY AND DOCUMENTS (GCC CLAUSE 10)	51
7.	INSURANCE (GCC CLAUSE 11)	52
8.	INCIDENTAL SERVICES (GCC CLAUSE 13)	52
9.	SPARE PARTS (GCC CLAUSE 14)	52
10.	WARRANTY (GCC CLAUSE 15)	52
11.	PAYMENT (GCC CLAUSE 16)	52
12.	PRICES (GCC CLAUSE 17)	52
13.	LIQUIDATED DAMAGES (GCC CLAUSE 23)	52
14.	RESOLUTION OF DISPUTES (GCC CLAUSE 28)	52
15.	GOVERNING LANGUAGE (GCC CLAUSE 29)	53
16.	APPLICABLE LAW (GCC CLAUSE 30)	53
17.	NOTICES (GCC CLAUSE 31)	53
18.	DUTIES AND TAXES (GCC CLAUSE-35)	53

## **Special Conditions of Contract**

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

### **1. Definitions (GCC Clause 1)**

GCC 1.1 (c) The Goods are: **Drugs / Medicines, Surgical Disposables, Medical Devices & Non-Drug Items (NDIs)**

GCC 1.1 (g) **The Procuring Agency is:** Director General Health Services, Khyber Pakhtunkhwa being the overall head of Government Medicine Coordination Cell (MCC) Health Department Government of Khyber Pakhtunkhwa; and

**The Purchasing Agency/ies include:** District Health Officers, Medical Superintendents and other Heads of the Primary, Secondary and / or Tertiary Level Health Care Institutions in the Health Department, Government of Khyber Pakhtunkhwa, including health related projects and / or vertical programs and / or interventions of / by the Health Department, Khyber Pakhtunkhwa and Health Facilities of the Prisons throughout Khyber Pakhtunkhwa.

GCC 1.1 (i) The Supplier is: "the individual or firm supplying the Goods and Services under this Contract" and includes the following:

- i) Manufacturer/s and / or Importer/s of drugs / medicines authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan, registered as such with the Drug Regulatory Authority of Pakistan (DRAP) for the quoted item/s falling under The Drug Act 1976 & Rules framed thereunder; and
- ii) **Manufacturer/s** of Medical Devices in Pakistan, registered as such with the DRAP for the quoted item/s and regulated under the DRAP Act 2012 and the Rules framed thereunder; and
- iii) **Importer/s** of Medical Devices, duly authorized by the goods' Principal Manufacturer or producer to import / supply the said goods in Pakistan, as registered and regulated as such for the quoted item/s under the DRAP Act 2012 and Rules framed thereunder; and
- iv) **Manufacturer/s** of Non-Drug Items (NDIs) in Pakistan; and
- v) **Importer/s** of NDIs, duly authorized by the goods' Principal Manufacturer or producer for import / supply of the said quoted goods in Pakistan.

### **Sample Provision:**

**GCC 1.1 (j)—The Project Site is: Director Drug Control, at Drug Cell of the Directorate General Health Services, Warsak Road, Old Fata Secretariat Peshawar.**

**GCC 8.1:** When required, the Focal Person of the bidder will be informed on phone or through email to provide samples of the items in sufficient / required quantity for examination / analysis / expert opinion to the office of Government MCC at bidder's own risk and cost at, and not later than, the time and date communicated. The samples will be non-returnable and no payment whatsoever shall be payable to bidder / Focal Person on this account in the name of price / transportation charges etc. or on the basis of any other context or reason or argument.

**2. Country of Origin (GCC Clause 3)**

All countries and territories as indicated in Part Two Section VI of the bidding documents, “Eligibility for the Provisions of Goods, Works, and Services in Government-Financed Procurement”.

**3. Performance Security (GCC Clause-7)**

**GCC 7.1—** The amount of performance security, as a percentage of the Contract Price, shall be: **Not Required.**

However, the bid security of Rs. 800,000/- from the successful bidders as received at the time of bids submission under GCC Clause 15, shall be retained by the Procuring Agency as Performance Security till the end of contract period and will be released back to successful bidders after the expiry of contract period, subject to the condition that all contractual obligations related to supplies are fulfilled. However, the warranty of the supplied goods, as issued by the Supplier under the clauses of contract agreement (Bid Form-6) and relevant applicable laws governing the nature of goods, e.g., the Drug Act 1976, The DRAP Act 2012 and rules framed there under shall remain in force and valid despite the discharge of Performance Security to the Supplier in accordance with GCC Clause-7 and 8.

**4. Standards (GCC Clause 4):** As mentioned in GCC clause 4.1.

**5. Inspections and Tests (GCC Clause 8 and in accordance with the clauses of contract with the Procuring Agency)**

- i. The Technical Evaluation shall be conducted by the Inspection Team/s of MCC expert/s constituted by the Technical and Evaluation (T&E) Committee and /or by the Selection and Rate Contracting Committee (S&RCC) of the Government MCC to:
  - a. undertake examination of the original documents as mentioned in the Bid Cover Sheet (Bid Form-1) of these SBDs, and the attested copies of which had been submitted by the bidder/s along with the technical bids; and
  - b. undertake the physical inspection of the relevant premises to verify the status of Current Good Manufacturing Practices (cGMP) Parameters for the quoted item/s as laid down in the Technical Evaluation Proformas (Section-V: Technical Specification of the Part-II of these SBDs); and
  - c. examine the original documents related to the fitness of the material of immediate container/s for storage and / or dispensing of the quoted drugs / medicines item/s, e.g., Certificate of Analysis, invoice, etc. of the material/s used in manufacturing of the immediate container of quoted drug / medicine item/s, including that of its stopper / lid / cap.
- ii. The bidder shall be disqualified for competition if Inspection Team/s declare that the bidder did not meet the mandatory requirements for qualification at the time of inspection as mentioned in the approved Technical Evaluation Proforma in these SBDs for various categories of Suppliers.
- iii. The technical and financial bid shall be in conformity to rule 39 (1) & (3) of the KPPRA Rules, any deviation from it, the bid shall be treated as non-responsive.
- iv. Medical Devices, Surgical Disposables and NDIs shall be examined and / or tested by MCC expert/s of the T&E Committee, and / or of the S&RCC of the Government MCC in a manner as deemed relevant and appropriate (including testing at Drug Testing Lab or elsewhere) for the purpose by the said expert/s, and as laid down, or

otherwise, in the applicable laws and Rules, for submission of technical report to the relevant forum/quarter for the needful.

- v. The samples of Medical Devices and Surgical Disposables shall be examined and tested for selected parameters by the Drug Testing Laboratory for submission of technical report/s to relevant forum/quarters for the needful.
- vi. To fulfill the relevant clauses of the contract agreement (Bid Form-6 of these SBDs) for testing of supplied goods, all the successful bidders for Drugs/Medicine, Surgical Disposables, Medical Devices falling under the Drugs Act 1976, before signing the Contract Agreement (Bid Form-6) shall provide to the Procuring Agency, the Testing Method/s and Lab. protocols to test their quoted item/s in the Drugs Testing Laboratory.
- vii. Any other appropriate method/arrangements may be adopted by the T&E Committee and / or S&RCC to assess and/or assure the quality of goods being purchased and / or supplied to the Procuring and / or Purchasing Agency/ies.
- viii. The application fee charges of @ Rs. 5000/bid seems rational to carry out the purpose of soliciting the bidding documents as the same is considered as fee not only considering the cost of the documents but to achieve multiple steps relating to the procurement process including the product wise evaluation of the firms, technical & performance evaluation of the disposable items at their premises across the country by the panels of Pharmacists, consultants (physicians, surgeons, etc.) and other experts/end users and quality assurance parameters / specifications through chemical analysis in adherence to the standard specification of the offer bid as per provision of The Drug Act and rules frame their under.

#### **6. Packing (GCC Clause 9)**

The successful bidder shall make supplies of quoted item/s in accordance with the following:

- i. Provisions contained in the GCC Clause 9 of these SBDs; and
- ii. Relevant clauses of contract agreement of Government MCC with the Supplier/s (Bid Form-6 of these SBDs – Rate Contract Agreement); and
- iii. In case of item/s falling in the category of drugs / medicines, the immediate container of drug / medicine shall comply with the official monograph requirements, as submitted by the bidder to the DRAP with the dossier at the time of registration of the said quoted item/s with the DRAP in accordance with applicable provisions contained in the prevailing laws and rules.

#### **7. Delivery and Documents (GCC Clause 10)**

**Applicable Delivery Mode:** Delivered Duty Paid (DDP) as per contract agreement of the successful bidder with the Procuring Agency.

The Supplier shall provide the following documents to the Purchasing Agency:

- i. Copies of the Supplier's invoice showing Goods' description, quantity, unit price, and total amount;
- ii. Usual transport documents which the buyer may require to take the goods;
- iii. Manufacturer's / Importer's prescribed warranty certificate;

The supplier shall be responsible to transport the item/s in a manner that the appropriate and required storage temperature is continuously and properly maintained during transportation from supplier till delivery to the Purchasing Agency/ies. In case of item/s requiring the maintenance of cold chain, the supplier shall be under obligation to provide valid and appropriate evidence to the Purchasing Agency to the effect that end

to end cold chain of the supplied item/s has adequately been maintained during transportation of the said item/s to the Purchasing Agency/ies.

**8. Insurance (GCC Clause 11)**

GCC 11.1— The Goods supplied under the Contract shall be delivered duty paid (DDP) under which risk is transferred to the buyer after having been delivered, hence insurance coverage is sellers' responsibility. Since the Insurance is seller's responsibility, they may arrange appropriate coverage.

**9. Incidental Services (GCC Clause 13) Not applicable.**

**10. Spare Parts (GCC Clause 14) Not Applicable.**

**11. Warranty (GCC Clause 15)**

For goods belonging to the categories of Drugs/Medicines, Medical Devices, Surgical Disposables and Cotton related materials, and falling under the Drugs Act 1976 and / or the DRAP Act-2012 and Rules framed thereunder, the Supplier, in addition to the terms and conditions of the Rate Contract Agreement with Procuring Agency (Bid Form-6), shall provide warranty to the Purchasing Agency under all the relevant Section/s of applicable government laws and rules.

In case of goods belonging to the categories of NDIs, the Supplier as per GCC Clause 15 and the clauses of Contract Agreement with the Procuring Agency (Bid Form-6), shall provide warranty to the Purchasing Agency for the duration as mentioned in GCC Clause-15 or till the expiry date of goods supplied, whichever is later.

**12. Payment (GCC Clause 16):**

GCC Clause 16 as well as under the terms and condition in Rate Contract Agreement (Bid Form-6) with the Procuring Agency.

Payment shall be made in **Pak. Rupees** in accordance with the relevant government rules, regulations and procedures.

**13. Prices (GCC Clause 17)**

- i) The bidder shall not quote price/s of any item/s which is/are higher than the prices quoted by the bidder across the country to any entity procuring the quoted item/s through public funding.
- ii) In case of Drugs/Medicines the bidder shall not quote the price more than the trade price of individual quoted item/s.
- iii) In case of Medical Devices, Surgical Disposables and NDIs, the bidder shall not quote the prices more than the prevailing market trade price of the quoted item/s for bulk purchases.

**14. Liquidated Damages (GCC Clause 23)**

As in relevant clauses of the Rate Contract Agreement signed by the Supplier with the Procuring Agency.

**15. Disputes Resolution (GCC Clause 28)**

The dispute resolution mechanism to be applied will be pursuant to relevant clauses of Rate Contract Agreement (Bid Form-6) between the Supplier and the Procuring Agency.

If at all required, the jurisdiction of Court shall be of Peshawar, Khyber Pakhtunkhwa.

**16. Governing Language (GCC Clause 29)**

The Governing Language shall be: **English.**

For various item/s related to drug / medicine category, the language of official Monograph of the quoted drug / medicine item/s, as registered with the DRAP, shall be acceptable for the bidding process.

**17. Applicable Law (GCC Clause 30)**

The Contract shall be interpreted in accordance with all the relevant laws of Islamic Republic of Pakistan which include, but not limited to, the following legislations:

- i. The KPPRA Act 2012
- ii. The KPPRA Rules 2014
- iii. The Drugs Act 1976 and Rules framed thereunder
- iv. The DRAP Act 2012 and Rules framed thereunder
- v. The General Financial Rules of the Government of Khyber Pakhtunkhwa and all the relevant laws, rules and regulations pertaining to budgeting and financial management of public funds.
- vi. The Employment of Children (ECA) Act 1991
- vii. The Bonded Labour System (Abolition) Act of 1992
- viii. The Factories Act 1934
- ix. The Contract Act 1872
- x. The companies Ordinance 1984, amended Companies Act 2017

**18. Notices (GCC Clause 31)**

GCC 31.1—Procuring Agency address for notice purposes:

**Office of the Director General Health Services**

Directorate General Health Services, Khyber Pakhtunkhwa,  
Warsak road, old FATA Secretariat Peshawar.

Tel: 091-9214084

091-9210269

Fax: 091-9210230

Email [mccdgdcp@gmail.com](mailto:mccdgdcp@gmail.com)

**Supplier's address for notice purposes:** As mentioned in their bidding documents

**19. Duties & Taxes (GCC clause 32)**

The Unit price quoted by the bidder shall be: **inclusive** of all applicable duties and taxes.

## Section IV. Schedule of Requirements (SOR)

### GOVERNMENT MEDICINE CO-ORDINATION CELL

### HEALTH DEPARTMENT GOVT. OF KHYBER PAKHTUNKHWA

### MCC FORMULARY FOR THE YEAR 2021-23

**NOTE:**

**1.**All powdered Injectables shall be supplied with sterile water for injection (Specified volume/quantity sufficient as per dossier of the product submitted by the manufacturer at the time of registration) in accordance to the **DRAP registered packing of drug.**

**2.**In case a bidder has been awarded marks during the technical evaluation for different parameters, the successful bidder(s) shall supply the said item/s with the quoted specification(s), against which the marks have been awarded, to the Purchasing Agency/ies throughout the validity period of the contract agreement.

**3.** For Narcotic analgesic drugs, i.e., Morphine, proper procedure and protocol of Government shall be followed by the Purchasing Agency/ies and Supplier/s.

#### AMOEBICIDES

S.No	Drug Name	Strength	Dosage form	volume
1	Metronidazole	200 mg	Tab.	
2	Metronidazole	400 mg	Tab.	
3	Metronidazole	200 mg/5ml	Susp.	60 ml
4	Metronidazole	500 mg	Inf.	100 ml
5	Metronidazole	0.75%	Vag. Gel	15gm
6	Metronidazole	0.75%	Vag. Gel	75gm
7	Nitazoxanide	500 mg	Tab.	
8	Nitazoxanide	100 mg/5ml	Susp.	30 ml
9	Tinidazole	300 mg	Tab.	

10	Tinidazole	500 mg	Tab.	
	<b>ANAESTHETIC &amp; ADJUVANT</b>			
11	Atracurium Besylate	10 mg/ml	Inj.	
12	Atracurium Besylate	10 mg/ml	Inj.	5 ml
13	Bupivacaine HCl	5 mg/ml	Inj.	10 ml
14	Bupivacaine Spinal	7.5 mg/ml	Inj.	2 ml
15	Cis-Atracurium	2mg/ml	Inj.	5 ml
16	Dexmedetomidine	0.1mg/ml	Inj.	2 ml
17	Glycopyrrolate + Neostigmine	0.5 mg+2.5mg	Inj.	1ml
18	Glycopyrrolate	0.2 mg/ml	Inj.	1ml
19	Halothane		Liq. for Inh.	250 ml
20	Isoflurane		Liq. for Inh.	100 ml
21	Ketamine HCl	50 mg/ml	Inj.	10 ml
22	Ketamine HCl	50 mg/ml	Inj.	2 ml
23	Lignocaine HCl	2%	Inj.	10 ml
24	Lignocaine HCl	4%	Topical Soln.	50 ml
25	Lignocaine HCl + Adrenaline	20mg/ml + 0.001% w/v	Inj.	10 ml
26	Lignocaine HCl + Adrenaline	1:80,000	Dental Ctg.	2 ml
27	Pancuronium	4mg/2ml	Inj.	2ml
28	Propofol	10 mg/ml	Inj.	20 ml
29	Rocuronium	10 mg/ml	Inj.	5 ml
30	Ropivacaine HCl	5mg/ ml	Inj.	10 ml

31	Sevoflurane		Liq. for Inh.	250 ml
32	Succinyl Choline	50 mg/ml	Inj.	2 ml
33	Thiopentone Sodium	500 mg/Vial	Inj. (Dry Powder)	
34	Vecuronium Bromide	4 mg/Ampule	Inj. (Dry powder)	
	<b>ANALGESICS, ANTI-INFLAMMATORY, ANTIPYRETICS DRUGS &amp; MUSCLE RELAXANTS</b>			
35	Aceclofenac	100 mg	Tab.	
36	Acetyl Salicylic Acid (Aspirin)	300 mg	Disper. Tab.	
37	Baclofen	10mg	Tab.	
38	Diclofenac Sodium	25 mg	Supp.	
39	Diclofenac Sodium	100 mg	Supp.	
40	Diclofenac Sodium (IM/IV for Infusion)	25 mg/ml	Inj.	3 ml
41	Diclofenac Sodium enteric coated	50 mg	Tab.	
42	Fentanyl Citrate	0.05mg/ml	Inj.	5 ml
43	Ibuprofen	200 mg	Tab.	
44	Ibuprofen	400 mg	Tab.	
45	Ibuprofen	200 mg/ 5 ml	Susp.	90 ml
46	Ibuprofen	100 mg/ 5ml	Susp.	90 ml
47	Ibuprofen	400ml	Inf.	100ml
48	Ketorolac	30 mg/ml	Inj.	1ml
49	Mefenamic Acid	250 mg	Tab.	
50	Mefenamic Acid	500 mg	Tab.	
51	Mefenamic Acid	50 mg/5ml	Susp.	60 ml

52	Meloxicam	15 mg	Tab.	
53	Meloxicam	7.5 mg	Tab.	
54	Morphine	15 mg	Inj.	
55	Morphine	10 mg	Cap.	
56	Morphine	30 mg	Cap.	
57	Nalbuphine	10 mg	Inj.	
58	Nalbuphine	20 mg	Inj.	
59	Paracetamol	80mg/0.8ml	Oral Drops	
60	Paracetamol	500 mg	Tab.	
61	Paracetamol	120 mg/ 5 ml	Susp.	60 ml
62	Paracetamol	250 mg/ 5ml	Susp.	60 ml
63	Paracetamol	150mg/ ml	Inj.	2 ml
64	Paracetamol	1000 mg	Inf.	100 ml
65	Paracetamol	150 mg	Supp.	
66	Paracetamol + Orphenadrine	450 mg/35 mg	Tab.	
67	Serratiopeptidase	5 mg	Tab.	
68	Tizanidine	4mg	Tab.	
69	Tramadol HCl	50 mg/ml	Inj.	2 ml
	<b>ANTHELMINTICS DRUGS</b>			
70	Albendazole	200 mg	Tab.	
71	Albendazole	100 mg/5ml	Susp.	10 ml
72	Levamisole	50 mg	Tab.	

73	Levamisole	150 mg	Tab.	
74	Levamisole	40 mg/5ml	Syp.	30 ml
75	Mebendazole	100 mg	Tab.	
76	Mebendazole	500 mg	Tab.	
77	Mebendazole	100 mg/5ml	Susp.	30 ml
78	Niclosamide	500 mg	Tab.	
79	Pyrantel pamoate	250 mg	Tab.	
	<b>ANTI NEOPLASTIC AGENTS / IMMUNOSUPPRESSANT/IMMUNO MODULATORY DRUGS</b>			
80	Azathioprine	50 mg	Tab.	
81	Basiliximab	20 mg/ vial	Inj.	
82	Bleomycin	15 mg	Inj.	
83	Chlorambucil	2 mg	Tab.	
84	Cyclophosphamide	500 mg/Vial	Inj.	
85	Cyclosporine-A	25 mg	Cap.	
86	Cyclosporine-A	50 mg	Cap.	
87	Cyclosporine-A	100 mg	Cap.	
88	Doxorubicin	10 mg/ Vial	Inj.	
89	Doxorubicin	50 mg/ Vial	Inj.	
90	Everolimus	5 mg	Tab.	
91	Everolimus	10 mg	Tab.	
92	Filgrastim	300 mcg	Inj.	

93	Hydroxy Urea	400 mg	Cap.	
94	Hydroxychloroquine	200 mg	Tab.	
95	Leflunomide	20 mg	Tab.	
96	Melphalan	2 mg	Tab.	
97	Melphalan	5 mg	Tab.	
98	Methotrexate	10 mg	Tab.	
99	Mitomycin	10 mg/ Vial	Inj.	
100	Mycophenolate Mofetil	250 mg	Tab. / Cap.	
101	Mycophenolate Mofetil	500 mg	Tab. / Cap.	
102	Mycophenolate Sodium	180 mg	Tab. / Cap.	
103	Mycophenolate Sodium	360 mg	Tab. / Cap.	
104	Sirolimus	1mg	Tab.	
105	Tacrolimus	1mg	Tab. /Cap.	
106	Tacrolimus	0.5 mg	Tab./ Cap.	
107	Tamoxifen	10 mg	Tab.	
108	Tamoxifen	20 mg	Tab.	
109	Thalidomide	100 mg	Tab. / Cap.	
110	Zoledronic Acid	4 mg /Vial	Inj.	

	<b>ANTIDOTES (DRUGS AND NON-DRUGS, e.g. ACTIVATED CHARCOAL)</b>			
111	Acetyl Cysteine		Inj.	
112	Activated Charcoal		Powder	
113	Activated Charcoal		Tab.	
114	Atropine Sulphate	1mg/ml	Inj.	1ml
115	Buprenorphine	0.3 mg/1 ml	Inj.	1 ml
116	Buprenorphine	2 mg	SL. Tab.	
117	Buprenorphine	8 mg	SL. Tab.	
118	Deferasirox	100 mg	Tab.	
119	Deferasirox	250 mg	Tab.	
120	Deferasirox	400 mg	Tab.	
121	Deferasirox	500 mg	Tab.	
122	Deferoxamine	500 mg	Inj.	
123	Dimercaprol	50 mg/ml	Inj.	
124	EDTA		Inj.	
125	Flumazenil	100 mcg/ml	Inj.	10 ml
126	Fomepizole	5 mg/ml	Inj.	
127	Glucagon	200 mg	Inj.	
128	Methylene Blue	10 mg/ml	Inj.	
129	N-acetylcysteine	200 mg	Sachet	

130	Naloxone HCl	0.4 mg / ml	Inj.	
131	Neostigmine	2.5 mg	Inj.	
132	Penicillamine	250 mg	Tab.	
133	Pralidoxime	20 mg/ml	Inj.	10 ml
134	Protamine Sulphate	10 mg/ml	Inj.	5 ml
135	Sodium Nitrite	30 mg	Inj.	
136	Sodium Thiosulfate	250 mg/ml	Inj.	
	<b>ANTI-FUNGAL DRUGS</b>			
137	Amphotericin-B	50 mg/Vial	Inj.	
138	Clotrimazole	1gm	Vag. Tab. with applicator	
139	Clotrimazole	1%	Vag. Cream with applicator	
140	Fluconazole	2 mg/ml	Inf.	50 ml
141	Fluconazole	50 mg	Tab. /Cap.	
142	Fluconazole	150 mg	Tab. /Cap.	
143	Fluconazole	50 mg/5 ml	Susp.	
144	Griseofulvin	500 mg	Tab.	
145	Griseofulvin	125 mg/5ml	Susp.	120 ml
146	Itraconazole	100 mg	Cap.	
147	Miconazole	2%	Skin Cream	10 gm

148	Miconazole	2%	Vag. Cream with Applicator	
149	Miconazole	2%	Oral Gel	
150	Nystatin	100,000 IU/5ml	Oral Drops	30 ml
151	Nystatin	100,000 IU	Vag. Tab. with applicator	
152	Terbinafine	250 mg	Tab.	
153	Voriconazole	200 mg	Inj.	
154	Voriconazole	200 mg	Tab.	
	<b>ANTI-HISTAMINES and ANTI-ALLERGIC DRUGS</b>			
155	Betahistine	8 mg	Tab.	
156	Betahistine	16 mg	Tab.	
157	Cetirizine	10 mg	Tab.	
158	Cetirizine	5 mg/5 ml	Syp.	60 ml
159	Chlorpheniramine Maleate	4 mg	Tab.	
160	Chlorpheniramine Maleate	2 mg/ 5 ml	Syp.	120 ml
161	Levocetirizine	2.5 mg/5 ml	Syp.	60 ml
162	Levocetirizine	5 mg	Tab.	
163	Loratadine	10 mg	Tab.	
164	Montelukast	10 mg	Tab.	
165	Montelukast	5 mg	Tab.	
166	Montelukast	4 mg	Sachet	

167	Pheniramine Maleate	25 mg/ml	Inj.	2 ml
	<b>ANTI-INFECTIVE DRUGS</b>			
168	Amikacin Sulphate	25 mg	Inj.	
169	Amikacin Sulphate	50 mg	Inj.	
170	Amikacin Sulphate	100 mg	Inj.	
171	Amikacin Sulphate	250 mg	Inj.	
172	Amikacin Sulphate	500 mg	Inj.	
173	Amoxycillin	250 mg	Cap.	
174	Amoxycillin	500 mg	Cap.	
175	Amoxycillin	125 mg/ 5ml	Dry Susp.	60 ml
176	Amoxycillin	125 mg/ 5ml	Dry Susp.	90 ml
177	Amoxycillin	500 mg/Vial	Inj.	
178	Amoxycillin	250mg /5ml	Dry Susp.	60 ml
179	Amoxycillin	250 mg /5ml	Dry Susp.	90 ml
180	Amoxicillin + Clavulanic Acid	250 mg/125mg (375mg)	Tab.	
181	Amoxicillin + Clavulanic Acid	500 mg/125mg (625 mg)	Tab.	
182	Amoxicillin + Clavulanic Acid	875 mg/125mg (1gm)	Tab.	
183	Amoxicillin + Clavulanic Acid	125 mg +31.5mg/5ml	Dry Susp.	90 ml
184	Amoxicillin + Clavulanic Acid	50 mg + 12.5mg/1ml	Oral Drops	20 ml

185	Amoxicillin + Clavulanic Acid	250 mg +62.5mg/5ml	Dry Susp.	90 ml
186	Amoxicillin + Clavulanic Acid	500 mg + 100mg/vial	Inj.	
187	Amoxicillin + Clavulanic Acid	1gm+200mg/V ial	Inj.	
188	Ampicillin	250 mg/Vial	Inj.	
189	Ampicillin	500 mg/Vial	Inj.	
190	Ampicillin + Cloxacillin	250 mg+ 250mg	Cap.	
191	Ampicillin + Cloxacillin	125mg +125mg/Vial	Inj.	
192	Ampicillin + Cloxacillin	250 mg + 250mg/vial	Inj.	
193	Ampicillin + Cloxacillin	125 mg + 125mg	Cap.	
194	Azithromycin	250 mg	Tab. / Cap.	
195	Azithromycin	500 mg	Tab. / Cap.	
196	Azithromycin	500 mg/Vial	Inj.	
197	Azithromycin	200 mg/5ml	Dry Susp.	15ml
198	Benzathine Penicillin	1.2 MIU/Vial	Inj.	
199	Benzyl Penicillin	10 Lac Units/Vial	Inj.	
200	Cefaclor	50mg / ml	Oral Drops	15 ml
201	Cefaclor	100mg/ml	Oral Drops	15 ml
202	Cefaclor	125mg/ 5ml	Susp.	60 ml
203	Cefaclor	250 mg /5ml	Susp.	60 ml

204	Cefazolin	500 mg/Vial	Inj.	
205	Cefazolin	1 gm/Vial	Inj.	
206	Cefepime	500 mg/vial	Inj.	
207	Cefepime	1 gm/vial	Inj.	
208	Cefixime	400 mg	Cap.	
209	Cefixime	100 mg/5ml	Dry Susp.	30ml
210	Cefixime	200 mg/5ml	Dry Susp.	30ml
211	Cefoperazone + Sulbactam	1gm/Vial	Inj.	
212	Cefoperazone + Sulbactam	2 gm/Vial	Inj.	
213	Cefotaxime Sodium	250 mg/Vial	Inj.	
214	Cefotaxime Sodium	500 mg/Vial	Inj.	
215	Cefotaxime Sodium	1gm/Vial	Inj.	
216	Cefpodoxime	100 mg	Tab.	
217	Cefpodoxime	40 mg/5ml	Dry Susp.	50 ml
218	Ceftaroline fosamil	600 mg/Vial	Inj.	
219	Ceftazidime	500 mg/Vial	Inj.	
220	Ceftazidime	1gm/Vial	Inj.	
221	Ceftriaxone	500 mg/Vial	Inj.	
222	Ceftriaxone	1gm/Vial	Inj.	

223	Ceftriaxone	2 gm Vial	Inj.	
224	Cefuroxime	1.5gm/Vial	Inj.	
225	Cefuroxime	250 mg	Tab.	
226	Cefuroxime	125 mg/5ml	Dry Susp.	
227	Cefuroxime	750 mg/Vial	Inj.	
228	Cephradine	250 mg	Cap.	
229	Cephradine	500 mg	Cap.	
230	Cephradine	1gm / Vial	Inj.	
231	Cephradine	500 mg / Vial	Inj.	
232	Cephradine	125mg / 5ml	Dry Susp.	90 ml
233	Cephradine	250 mg / 5ml	Dry Susp.	90 ml
234	Ciprofloxacin	250 mg	Tab.	
235	Ciprofloxacin	500 mg	Tab.	
236	Ciprofloxacin	200 mg/100ml	Inf.	100 ml
237	Ciprofloxacin	400 mg/100ml	Inf.	100 ml
238	Clarithromycin	250 mg	Tab.	
239	Clarithromycin	500 mg	Tab.	
240	Clarithromycin	250 mg/5ml	Dry Susp.	60 ml
241	Clarithromycin	125 mg/5ml	Dry Susp.	60 ml

242	Clarithromycin	125 mg/ 5 ml	Dry powder oral drops	25 ml
243	Clarithromycin	500 mg/Vial	Inj.	
244	Clindamycin	150 mg/ml	Inj.	2 ml
245	Cloxacillin	250 mg /Vial	Inj.	
246	Cloxacillin	250 mg	Cap.	
247	Colistimethate Sodium	2 MIU/vial	Inj.	
248	Colistimethate Sodium	1 MIU/vial	Inj.	
249	Co-Trimoxazole (Sulphamethoxazole +Trimethoprim)	400 mg + 80mg	Tab.	
250	Co-Trimoxazole (Sulphamethoxazole +Trimethoprim)	800 mg + 160mg	Tab.	
251	Co-Trimoxazole (Sulphamethoxazole +Trimethoprim)	400 mg + 80mg/5 ml	Susp.	50ml
252	Co-Trimoxazole (Sulphamethoxazole +Trimethoprim)	200mg + 40mg/5ml	Susp.	50ml
253	Dapsone	25 mg	Tab.	
254	Dapsone	100 mg	Tab.	
255	Doxycycline	100 mg	Cap.	
256	Ethambutol	400mg	Tab.	
257	Ethambutol	100mg	Disper. Tab.	
258	Flucloxacillin + Amoxicillin	250 mg + 250mg/ Vial	Inj.	

259	Flucloxacillin + Amoxicillin	250 mg + 250mg	Cap.	
260	Fosfomycin	500 mg	Cap.	
261	Fosfomycin	3 gm	Sachet	
262	Gentamicin Sulphate	20 mg/ml	Inj.	1ml
263	Gentamicin Sulphate	40 mg/ml	Inj.	2 ml
264	Imipenem + Cilastatin	500 mg+500mg / Vial	Inj.	
265	Isoniazid	300mg	Tab.	
266	Isoniazid	100mg	Disper. Tab.	
267	Levofloxacin	5 mg/ml	Inf.	100 ml
268	Levofloxacin	250 mg	Tab.	
269	Levofloxacin	500 mg	Tab.	
270	Lincomycin	500 mg	Cap.	
271	Lincomycin	300 mg/ml	Inj.	2 ml
272	Linezolid	600mg	Tab.	
273	Linezolid	2 mg/ml	Inf.	100 ml
274	Linezolid	2 mg/ml	Inf.	300 ml
275	Meropenem	500 mg/Vial	Inj.	
276	Meropenem	1gm /Vial	Inj.	
277	Minocycline	100 mg	Tab.	

278	Moxifloxacin	400 mg	Tab.	
279	Moxifloxacin	400 mg/250ml	Inf.	250 ml
280	Nitrofurantoin	100 mg	Tab.	
281	Oxytetracycline	250mg	Cap.	
282	Piperacillin +Tazobactam	2 gm+0.25gm (2.25gm)/Vial	Inj.	
283	Piperacillin +Tazobactam	4 g/0.5 g (4.5gm)/Vial	Inj.	
284	Pyrazinamide	400mg	Tab.	
285	Rifampicin	150 mg	Tab. / Cap.	
286	Rifampicin	300 mg	Tab. / Cap.	
287	Rifampicin	450 mg	Tab. / Cap.	
288	Rifampicin	600 mg	Tab. / Cap.	
289	Rifampicin	100 mg/5ml	Susp.	60 ml
290	Rifampicin +Isoniazid + Pyrazinamide + Ethambutol	150mg+75mg+ 400mg+275mg	Tab.	
291	Rifampicin+ Isoniazid+ Pyrazinamide	75mg + 50mg+150mg	Disper. Tab.	
292	Rifampicin +Isoniazid	150mg + 75mg	Tab.	
293	Rifampicin+ Isoniazid	75mg+50mg	Disper. Tab.	
294	Rifaximin	200 mg	Tab.	
295	Rifaximin	550 mg	Tab.	
296	Streptomycin Sulphate	1gm/Vial	Inj.	

297	Tigecycline	50 mg /Vial	Inj.	
298	Vancomycin	500 mg/Vial	Inj.	
299	Vancomycin	1gm/Vial	Inj.	
	<b>ANTI-MALARIAL DRUGS</b>			
300	Amodiaquine	150 mg/5 ml	Susp.	20 ml
301	Amodiaquine	150 mg	Tab.	
302	Artemether	80 mg/ml	Inj.	1ml
303	Artemether + Lumefantrine	40 mg/240mg	Tab.	
304	Artemether + Lumefantrine	80 mg/480mg	Tab.	
305	Artemether + Lumefantrine	15 mg/ 90 mg/5ml	Susp.	60ml
306	Artesunate	60 mg/Vial	Inj.	
307	Artesunate	120 mg/Vial	Inj.	
308	Artesunate + Sulfadoxine + Pyrimethamine	100mg+500mg+25 mg	Tab. Co- Blister	
309	Artesunate + Sulfadoxine + Pyrimethamine	50mg+500mg+25 mg	Tab. Co- Blister	
310	Chloroquine Phosphate	250 mg	Tab.	
311	Chloroquine Phosphate	50 mg/5ml	Syp.	60 ml
312	Dihydro artemisinin + Piperaquine Phosphate	15 mg + 120mg	Sachet	
313	Dihydroartemisinin+ Piperaquine Phosphate	40 mg + 320mg	Tab./ Cap.	
314	Primaquine	7.5 mg	Tab.	

315	Primaquine	15mg	Tab.	
316	Pyrimethamine	25 mg	Tab.	
317	Quinine Dihydrochloride	300 mg	Tab.	
318	Quinine Dihydrochloride	300 mg/ml	Inj.	2 ml
319	Sulfadoxine + Pyrimethamine	501 mg + 25mg	Tab.	
320	Sulfadoxine + Pyrimethamine	500 mg + 25mg/5ml	Susp.	15 ml
	<b>ANTI-VIRAL DRUGS</b>			
321	Acyclovir	200 mg	Tab.	
322	Acyclovir	250 mg/Vial	Inj.	
323	Acyclovir	500 mg/Vial	Inj.	
324	Famciclovir	250 mg	Tab.	
325	Ganciclovir	250 mg	Cap.	
326	Ganciclovir	500 mg/Vial	Inj.	
327	Oseltamivir	75mg	Cap.	
	<b>BLOOD FORMING DRUGS, COAGULANTS, ANTICOAGULANTS &amp; ANTI-ANAEMIC</b>			
328	Alteplase	2 mg	Inj.	
329	Alteplase	50 mg	Inj.	
330	Alteplase	100 mg	Inj.	
331	Enoxaparin	20 mg	Inj.	0.2 ml

332	Enoxaparin	40 mg	Inj.	0.4 ml
333	Enoxaparin	60 mg	Inj.	0.6 ml
334	Enoxaparin	80 mg	Inj.	0.8 ml
335	Epoetin- $\alpha$	2000 IU/Vial	Inj.	
336	Epoetin- $\alpha$	4000 IU /Vial	Inj.	
337	Epoetin- $\alpha$	10,000 IU/Vial	Inj.	
338	Epoetin- $\beta$	2000 IU/Vial	Inj.	
339	Epoetin- $\beta$	5000 IU/Vial	Inj.	
340	Epoetin- $\beta$	10,000 IU/Vial	Inj.	
341	Fondaparinux Sodium	2.5 mg	Inj.	
342	Fondaparinux Sodium	7.5 mg	Inj.	
343	Factor IX	500 IU/Vial	Inj.	
344	Factor VII	1mg /Vial	Inj.	
345	Factor VII	5mg /Vial	Inj.	
346	Factor VIII	250 IU/vial	Inj.	
347	Ferrous Fumarate + Folic Acid	185mg + 400mcg	Tab.	
348	Ferrous Sulphate	200 mg	Tab.	
349	Ferrous Sulphate	100 mg/5ml	Syp.	120 ml
350	Folic Acid	5 mg	Tab.	

351	Heparin Sodium	5000 IU/ml	Inj.	5ml
352	Iron Hydroxide poly maltose complex	100 mg	Tab.	
353	Iron Hydroxide poly maltose complex	50 mg/5ml	Syp	60 ml
354	Iron Hydroxide poly maltose complex	50 mg/ml	Oral Drops	30 ml
355	Iron Isomaltoside	100 mg	Inj.	1ml
356	Iron Sucrose	20 mg/ml	Inj.	5 ml
357	Mecobalamin	500 mcg	Inj.	1ml
358	Methoxy PEG Epoetin-β	50 mcg	Inj.	0.3 ml
359	Methoxy PEG Epoetin-β	75 mcg	Inj.	0.3 ml
360	Methoxy PEG Epoetin-β	100 mcg	Inj.	0.3 ml
361	Methoxy PEG Epoetin-β	150 mcg	Inj.	0.3 ml
362	Methoxy PEG Epoetin-β	200 mcg	Inj.	0.3 ml
363	Phytomenadione Inj. (vit-K1)	2mg/ml	Inj.	1ml
364	Rivaroxaban	10 mg	Tab.	
365	Rivaroxaban	15 mg	Tab.	
366	Rivaroxaban	20 mg	Tab.	
367	Tranexamic Acid	500 mg	Cap.	
368	Tranexamic Acid	250 mg	Inj.	5 ml
369	Tranexamic Acid	500 mg	Inj.	5 ml

370	Warfarin Sodium	1 mg	Tab.	
371	Warfarin Sodium	2.5 mg	Tab.	
372	Warfarin Sodium	5 mg	Tab.	
	<b>CARDIOVASCULAR AND DIURETIC DRUGS</b>			
373	Acetazolamide.	250 mg	Tab.	
374	Acetyl Salicylic Acid (Aspirin) EC.	75 mg	Tab.	
375	Adenosine		Inj.	
376	Adrenaline	1mg/ml	Inj.	1ml
377	Amiodarone HCl	200 mg	Tab.	
378	Amiodarone HCl	100 mg	Tab.	
379	Amiodarone HCl	150 mg/ml	Inj.	3 ml
380	Amlodipine Besylate	5 mg	Tab.	
381	Amlodipine Besylate	10 mg	Tab.	
382	Amlodipine+Valsartan	5mg/80 mg	Tab.	
383	Amlodipine+Valsartan	5mg/160 mg	Tab.	
384	Amlodipine+Valsartan	10 mg/160 mg	Tab.	
385	Atenolol	50 mg	Tab.	
386	Atenolol	100 mg	Tab.	
387	Bisoprolol	2.5mg	Tab.	
388	Bisoprolol	5 mg	Tab.	

389	Bisoprolol	10 mg	Tab.	
390	Bosenton	62.5mg	Tab.	
391	Candesartan	4 mg	Tab.	
392	Candesartan	8 mg	Tab.	
393	Candesartan	16 mg	Tab.	
394	Candesartan + Hydrochlorothiazide	16 mg+12.5mg	Tab.	
395	Captopril	25 mg	Tab.	
396	Carvedilol	6.25 mg	Tab.	
397	Carvedilol	12.5 mg	Tab.	
398	Carvedilol	25 mg	Tab.	
399	Clopidogrel	75 mg	Tab.	
400	Clopidogrel	300 mg	Tab.	
401	Digoxin	500 mcg (0.5mg)	Inj.	2ml
402	Digoxin	250 mcg	Tab.	
403	Digoxin	50 mcg/ml	Oral Soln.	
404	Dobutamine HCl	50 mg/ml	Inj.	5 ml
405	Dopamine HCl	40 mg/ml	Inj.	5 ml
406	Dopamine HCl	80 mg/ml	Inj.	10 ml
407	Furosemide	20 mg	Tab.	
408	Furosemide	40 mg	Tab.	
409	Furosemide	10 mg/ml	Inj.	2ml

410	Glyceryl Trinitrate	0.5 mg	SL. Tab.	
411	Glyceryl Trinitrate	2.6 mg	Tab.	
412	Glyceryl Trinitrate	6.4 mg	Tab.	
413	Glyceryl Trinitrate	5 mg	Patch	
414	Glyceryl Trinitrate	400 mcg	Buccal Spray	200 doses
415	Hydralazine	20 mg	Inj.	
416	Hydralazine	25 mg	Tab.	
417	Hydralazine	50 mg	Tab.	
418	Hydrochlorothiazide	25 mg	Tab.	
419	Isoprenaline	1 mg/ml	Inj.	2 ml
420	Isosorbide Dinitrate	1mg/ml	Inj.	10 ml
421	Isosorbide Dinitrate	5 mg	Tab.	
422	Isosorbide Dinitrate	10 mg	Tab.	
423	Isosorbide-5-Mononitrate	20 mg	Tab.	
424	Isosorbide-5-Mononitrate	40 mg	Tab.	
425	Labetalol	50 mg	Inj.	10 ml
426	Lisinopril	5 mg	Tab.	

427	Lisinopril	10 mg	Tab.	
428	Losartan + Hydrochlorothiazide	50 mg+12.5mg	Tab.	
429	Losartan Potassium	25 mg	Tab.	
430	Losartan Potassium	50 mg	Tab.	
431	Methyldopa	250 mg	Tab.	
432	Methyldopa	250 mg	Inj.	
433	Metoprolol	25 mg	Tab.	
434	Metoprolol	50 mg	Tab.	
435	Metoprolol	100 mg	Tab.	
436	Metoprolol	1mg/ml	Inj.	5 ml
437	Metolazone	5 mg	Tab.	
438	Milrinone	1mg/ml	Inj.	10ml
439	Nifedipine	10 mg	Cap.	
440	Nifedipine	30 mg	ER-Tab.	
441	Nifedipine	30mg	Tab.	
442	Nitroglycerin	1mg/ml	Inj.	
443	Noradrenaline / Norepinephrine	1mg/ml	Inj.	4 ml
444	Phenylephrine	10 mg	Inj.	
445	Procaine + Magnesium chloride+ Potassium chloride	0.27 mg/10ml+ 3.25mg/10ml + 1.19mg/10ml	Inj.	10 ml

446	Propranolol	10 mg	Tab.	
447	Propranolol	40 mg	Tab.	
448	Ramipril	5 mg	Tab.	
449	Rosuvastatin	10 mg	Tab.	
450	Sodium Nitroprusside	25mg/ml	Inj.	2ml
451	Spironolactone	100 mg	Tab.	
452	Streptokinase	1.5 MIU/vial	Inj.	
453	Valsartan	40 mg	Tab.	
454	Valsartan	80 mg	Tab.	
455	Valsartan + Hydrochlorothiazide	80 mg+12.5mg	Tab.	
456	Valsartan + Sacubitril	51mg + 49mg	Tab.	
457	Verapamil	40 mg	Tab.	
458	Verapamil	80 mg	Tab.	
459	Verapamil	2.5 mg/ml	Inj.	2 ml
	<b>EAR, NOSE AND THROAT PREPARATIONS</b>			
460	Betamethasone	0.10%	Ear /Nasal Drops	7.5 ml
461	Betamethasone + Neomycin	0.1% + 0.5%	Ear/Nasal Drops	7.5 ml
462	Ciprofloxacin HCl	0.30%	Ear Drops	5 ml
463	Fluticasone	50 mcg/Actu.	Nasal Spray	15ml

464	Lignocaine + Polymyxin	50mg/ml+10,000 IU/ml	Ear Drops	5ml
465	Soda Glycerin (Sodium Bicarbonate + Glycerin)	5% +30%	Ear Drops	10 ml
466	Sodium Chloride	0.65 % w/v	Nasal Drops	30 ml
467	Xylometazoline HCl	0.05%	Nasal Drops	15ml
468	Xylometazoline HCl	0.10%	Nasal Spray	15ml
<b>GASTROINTESTINAL DRUGS</b>				
469	Aluminum Hydroxide + Magnesium Hydroxide + Simethicone		Susp.	
470	Bacillus Clausii Spores	2 Billion/ 5ml	Susp.	
471	Bisacodyl	5 mg	Tab.	
472	Dimenhydrinate	12.5mg/4ml	Syp.	60 ml
473	Dimenhydrinate	50 mg/ml	Inj.	1 ml
474	Dimenhydrinate	50 mg	Tab.	
475	Domperidone	10 mg	Tab.	
476	Domperidone	5 mg/5ml	Susp.	120 ml
477	Drotaverine	40 mg	Tab.	
478	Drotaverine	20 mg/ml	Inj.	2ml
479	Famotidine	40 mg	Tab.	

480	Glycerin Suppositories		Supp.	
481	Lactulose	3.35gm/5ml	Syp.	120ml
482	Loperamide	2mg	Cap.	
483	Metoclopramide HCl	5mg/ml	Inj.	2ml
484	Octreotide Acetate	0.1mg/ml	Inj.	1ml
485	Omeprazole	40 mg / Vial	Inj.	
486	Ondansetron	8 mg	Tab.	
487	Ondansetron	2 mg/ml	Inj.	4 ml
488	Pantoprazole	20 mg	Tab.	
489	Phloroglucinol + Trimethyl Phloroglucinol	80 mg + 80 mg	Tab.	
490	Phloroglucinol + Trimethyl Phloroglucinol	40 mg + 0.04mg	Inj.	4 ml
491	Prucalopride	2 mg	Tab.	
492	Simethicone	40 mg/ml	Oral Drops	30 ml
493	Sodium Phosphate + Sodium Bi-Phosphate	7.2 gm + 19.2gm	Enema	120ml
494	Sodium Citrate + Sodium Lauryl Sulphate + Glycerin	450mg+75mg+ 90%	Enema	10ml
495	Sodium Bicarbonate + Peppermint		Tab.	
496	Terlipressin	1mg / Vial	Inj.	
497	Zinc Sulphate	20 mg	Tab.	
498	Zinc Sulphate	20 mg/5ml	Syp.	60 ml
	<b>HORMONES &amp; DRUGS ACTING ON ENDOCRINE SYSTEM</b>			

499	Carbimazole	5 mg	Tab.	
500	Clomiphene Citrate	50 mg	Tab.	
501	Dexamethasone	0.5 mg	Tab.	
502	Dexamethasone	4 mg/ml	Inj.	1ml
503	Dinoprostone	3 mg	Vag. Tab.	
504	Empagliflozin	10 mg	Tab.	
505	Empagliflozin	25 mg	Tab.	
506	Fludrocortisone	0.1 mg	Tab.	
507	Glibenclamide	5 mg	Tab.	
508	Gliclazide	80 mg	Tab.	
509	Glimepiride	1mg	Tab.	
510	Glimepiride	2mg	Tab.	
511	Glimepiride	3mg	Tab.	
512	Glimepiride	4mg	Tab.	
513	Glimepiride + Metformin	1 mg/500mg	Tab.	
514	Glimepiride + Metformin	2 mg/500mg	Tab.	
515	Human chorionic gonadotropin	1500 IU	Inj.	
516	Human chorionic gonadotropin	5000 IU	Inj.	
517	Hydrocortisone	100 mg/Vial	Inj.	
518	Hydrocortisone	250 mg/Vial	Inj.	
519	Hydroxy progesterone	250mg/ml	Inj.	1 ml
520	Insulin 70/30 Premixed (Human)	100 IU /ml	Inj.	10ml

521	Insulin Regular (Human)	100 IU/ml	Inj.	10ml
522	Insulin Glargine	100 IU/ml	Inj.	10ml
523	Insulin Lispro	100 IU/ml	Inj.	10ml
524	Insulin Isophane	100 IU/ml	Inj.	10ml
525	Mestranol + Norethisterone	50 mcg + 1 mg	Tab.	
526	Metformin HCl	500mg.	Tab.	
527	Methyl Prednisolone	500mg Vial	Inj.	
528	Methyl Prednisolone	1gm Vial	Inj.	
529	Methylergometrine Maleate	0.2 mg/ml	Inj.	1 ml
530	Misoprostol	200 mcg	Tab.	
531	Oxybutynin	5mg	Tab.	
532	Oxytocin	5 IU/ml	Inj.	1 ml
533	Oxytocin	10 IU/ml	Inj.	1 ml
534	Prednisolone	5 mg	Tab.	
535	Propylthiouracil	50 mg	Tab.	
536	Prostaglandin F2	5mg/ml	Inj.	1ml
537	Sitagliptin + Metformin	50 mg/500 mg	Tab.	
538	Sitagliptin + Metformin	50mg /1000 mg	Tab.	
539	Thyroxin Sodium	50 mcg	Tab.	
540	Triamcinolone Acetonide	40 mg	Inj.	1 ml
541	Vildagliptin	50 mg	Tab.	
	<b>IMMUNOLOGICAL / BIOLOGICAL DRUGS</b>			

542	Anti Gas Gangrene Serum	30000 Units	Inj.	
543	Anti-Rabies Serum	200 IU/ml		5 ml
544	Anti-Tetanus Serum	1500 IU	Inj.	1ml
545	Anti-Tetanus Serum	10,000 IU	Inj.	
546	Anti-thymocyte globulin (ATG)		Inj.	
547	Cholera Vaccine		Inj.	
548	Diphtheria Anti-Toxin	20,000 IU	Inj.	
549	Diphtheria Anti-Toxin	10,000 IU	Inj.	
550	Hepatitis B Immunoglobulin (Adult)		Inj.	
551	Hepatitis B Immunoglobulin (Neonatal)		Inj.	
552	Human Immunoglobulins for IV administration	5%	Inj.	
553	Human Immunoglobulins for IV administration	10%	Inj.	
554	Human Diploid Cell Rabies Vaccine (HDCV)		Inj.	
555	Meningococcal Vaccine (WHO Prequalified)		Inj.	
556	Mumps Measles Rubella Vaccine (MMR)		Inj.	
557	Mumps Vaccine		Inj.	
558	Pneumococcal (WHO Prequalified)	PCV13	Inj.	
559	Pneumococcal (WHO Prequalified)	PPSV23	Inj.	
560	Purified Chick Embryo Cell Rabies Vaccine (PCECV)		Inj.	

561	Purified Vero Cell Rabies Vaccine (PVRV)		Inj.	
562	Primary Hamster Kidney Cell Rabies vaccine (PHKCV)		Inj.	
563	Purified Duck Embryo Rabies vaccine (PDEV)		Inj.	
564	Rabies Immunoglobulin (Human)	150 IU/ml	Inj.	
565	Rho (D) Immune globulin	300 mcg	Inj.	
566	Rituximab	500 mg	Inj.	50ml
567	Secukinumab	150 mg	Inj.	
568	Scorpion Venom Antiserum		Inj.	
569	Snake Venom Antiserum		Inj.	
570	Tetanus Immunoglobulin (Human)	250 IU	Inj.	
571	Tetanus Toxoid	0.5 ml	Inj.	
572	Trivalent Influenza Vaccine (WHO Prequalified)		Inj.	
573	Typhoid Vaccine		Inj.	
	<b>INTRAVENOUS FLUIDS, ELECTROLYTES AND PARENTERAL NUTRITION</b>			
574	Amino Acids Solutions	3%, 4%, 7%, 8%, 5%, 10% & 20%	I/V Inf.	500 ml
575	Balanced electrolyte solution		I/V Inf.	500 ml
576	Calcium Chloride		Inj.	

577	Calcium Chloride, Glucose, Potassium Chloride, Sodium Acetate	0.2g/L, 5% w/v, 1.5g/L, 3.13g/L	I/V Inf.	500ml
578	Calcium Chloride, Glucose, Potassium Chloride, Sodium Acetate	0.2g/L, 5% w/v, 1.5g/L, 3.13g/L	I/V Inf.	1000ml
579	Calcium Gluconate		Inj.	10ml
580	Dextrose Soln.	0.25	I/V Inf.	25ml
581	Dextrose Soln.	0.25	I/V Inf.	1000ml
582	Dextrose Soln.	0.05	I/V Inf.	100ml
583	Dextrose Soln.	0.05	I/V Inf.	500ml
584	Dextrose Soln.	0.05	I/V Inf.	1000ml
585	Dextrose + Sodium Chloride	5% + 0.45%	I/V Inf.	500ml
586	Dextrose + Sodium Chloride	5% + 0.9%	I/V Inf.	500ml
587	Dextrose + Sodium Chloride	5% + 0.9%	I/V Infusion.	1000ml
588	Flavored Oral Re-hydration Salt WHO approved formula.	Sodium Chloride (3.5 g/L), Glucose Anhydrous (20g/L) Potassium Chloride (1.5g/L) Trisodium Citrate (2.9g/L)	Sachet	

589	Flavored Oral Rehydration Salt (Low Osmolarity)	Sodium Chloride Sachet (2.6 g/L) Glucose Anhydrous (13.5 g/L) Potassium Chloride (1.5 g/L) Trisodium citrate (2.9 g/L)	Sachet	
590	Gelatin Polypeptide	0.035	I/V Inf.	500 ml
591	Gelatin Polypeptide	0.04	I/V Inf.	500 ml
592	Glycine Irrigation solution		I/V Inf.	4000 ml
593	Hemodialysis Solution		Soln.	4000 ml
594	Lipid Emulsion	0.2	I/V Inf.	250 ml
595	Magnesium Sulphate	500 mg/ml	Inj.	2ml
596	Magnesium Sulphate	500 mg/ml	Inj.	10 ml
597	Mannitol	0.2	I/V Inf.	500 ml
598	Normal Saline	0.009	I/V Inf.	100 ml
599	Normal Saline	0.009	I/V Inf.	500 ml
600	Normal Saline	0.009	I/V Inf.	1000 ml
601	Peritoneal Dialysis Soln.		Soln	1000 ml
602	Peritoneal Dialysis Soln.		Soln	2000 ml
603	Peritoneal Dialysis Soln.		Soln	4000 ml
604	Potassium Chloride	1 gm/ 5ml	Syp.	120 ml

605	Potassium Chloride	7.46% w/v	Inj.	25ml
606	Potassium Chloride	500 mg	SR-Tab.	
607	Ringer's Lactate + Dextrose 5% Soln.		I/V Inf.	500 ml
608	Ringer's Lactate + Dextrose 5% Soln.		I/V Inf.	1000 ml
609	Ringer's Lactate Soln.		I/V Inf.	500 ml
610	Ringer's Lactate Soln.		I/V Inf.	1000 ml
611	Salt free Albumin	20% Soln.	I/V Inf.	50 ml
612	Salt free Albumin	20% Soln.	I/V Inf.	100 ml
613	Sodium Acid Citrate	1.315 gm/ 5 ml	Liq.	120 ml
614	Sodium Bicarbonate	0.084	I/V Soln.	
615	Sodium Chloride + Dextrose	0.18 % + 4.3%	I/V Inf.	500ml
616	Sterile Water for Injection	5 ml	Inj.	
617	Total Parenteral Nutrition (Glucose, Sodium Phosphate, Zinc)		IV Inf.	1250 ml
	<b>MISCELLANEOUS THERAPEUTICS</b>			
618	Allopurinol	100 mg	Tab.	
619	Allopurinol	300 mg	Tab.	
620	Beractant	25mg/ml	Inj.	
621	Cinacalcet HCL	30 mg	Tab.	

622	Febuxostat	40 mg	Tab.	
623	Febuxostat	80 mg	Tab.	
624	Hyaluronic Acid		Inj.	
625	Ibandronic Acid	1mg/ml	Inj.	3 ml
626	Ibandronic Acid	150mg	Tab.	
627	Liquid Paraffin			450 ml
628	Proactant alfa	120 mg/ 1.5 ml	Inj.	
629	Proactant alfa	240 mg/ 3 ml	Inj.	
630	Sevelamer Carbonate	400mg	Tab.	
631	Sodium tetradecyl sulphate	10mg/ ml (1%)	Inj.	2ml
632	Sodium tetradecyl sulphate	30mg/ml (3%)	Inj.	2 ml
633	Solifenacin Succinate	10mg	Tab.	
634	Tamsulosin HCl	0.4mg	Cap.	
635	Tamsulosin HCl + Dutasteride	0.4 mg+ 0.5mg	Cap.	
	<b>PSYCHOTROPIC AND ANTICONVULSANT DRUGS</b>			
636	Alprazolam	0.25 mg	Tab.	
637	Alprazolam	0.5 mg	Tab.	
638	Amitriptyline HCl	25 mg	Tab.	
639	Aripiprazole	15 mg	Tab.	

640	Carbamazepine	200 mg	Tab.	
641	Carbamazepine	100 mg / 5 ml	Syp.	120 ml
642	Chlorpromazine HCl	100 mg	Tab.	
643	Citalopram	10 mg	Tab.	
644	Citicoline	125 mg/ml	Inj.	2 ml
645	Citicoline	250 mg/ml	Inj.	2 ml
646	Clomipramine HCl	25 mg	Tab.	
647	Clonazepam	0.5 mg	Tab.	
648	Clonazepam	2 mg	Tab.	
649	Clonazepam	0.25% w/v	Oral Drops	10 ml
650	Clozapine	25mg	Tab.	
651	Clozapine	100 mg	Tab.	
652	Co-dergocrine mesylate	1.5 mg	Tab.	
653	Desvenlafaxine	50 mg	Tab.	
654	Desvenlafaxine	100 mg	Tab.	
655	Diazepam	10 mg/ml	Inj.	2 ml
656	Duloxetine	30 mg	Tab.	
657	Duloxetine	60 mg	Tab.	
658	Divalproex Sodium	250 mg	Tab.	

659	Divalproex Sodium	500 mg	Tab.	
660	Dothiepin HCl (Dosulepin HCl)	25mg	Tab.	
661	Dothiepin HCl (Dosulepin HCl)	75 mg	Tab.	
662	Escitalopram	10 mg	Tab.	
663	Fluoxetine HCl	20 mg	Cap.	
664	Flupenthixol	40 mg/ml	Inj.	2 ml
665	Fluphenazine Decanoate	25 mg/ml	Inj.	1 ml
666	Haloperidol	2 mg/ ml	Oral Drops	15 ml
667	Haloperidol	5 mg	Tab.	
668	Haloperidol	5 mg	Inj.	1 ml
669	Imipramine	25 mg	Tab.	
670	Lamotrigine	50 mg	Tab.	
671	Levodopa + Carbidopa	250 mg+25mg	Tab.	
672	Levetiracetam	250 mg	Tab.	
673	Levetiracetam	500mg	Tab.	

674	Levetiracetam	100 mg/ml	Inj.	5 ml
675	Lithium Carbonate	400 mg	Tab.	
676	Midazolam	1 mg/ml	Inj.	5 ml
677	Mirtazapine	15mg	Tab.	
678	Olanzapine	5mg	Tab.	
679	Olanzapine	10 mg	Tab.	
680	Oxcarbazepine	300 mg	Tab.	
681	Oxcarbazepine	600 mg	Tab.	
682	Phenobarbital	30 mg	Tab.	
683	Phenobarbital	200 mg	Inj.	1ml
684	Phenobarbital	20 mg/5ml	Elixir	60 ml
685	Phenytoin Sodium	100 mg	Tab. /Cap.	
686	Phenytoin Sodium	30 mg/5 ml	Susp.	
687	Phenytoin Sodium		Inj.	
688	Piracetam	200 mg/ml	Inj.	5ml

689	Pregabalin	50 mg	Cap.	
690	Pregabalin	75mg	Cap.	
691	Pregabalin	150 mg	Cap.	
692	Prochlorperazine Maleate	5 mg	Tab.	
693	Prochlorperazine Maleate	12.5 mg.	Inj.	1 ml
694	Procyclidine HCl	5mg	Tab.	
695	Procyclidine HCl	5 mg/ml	Inj.	2 ml
696	Quetiapine	100 mg	Tab.	
697	Risperidone	2mg	Tab.	
698	Risperidone	4 mg	Tab.	
699	Selegiline	5 mg	Tab	
700	Sertraline	100 mg	Tab.	
701	Sodium Valproate	250 mg/5ml	Syp.	120 ml
702	Topiramate	50 mg	Tab.	
703	Trifluoperazine	5 mg	Tab.	

704	Valproate Sodium	500 mg/5ml	Inj.	
705	Valproate Sodium	500 mg/5ml	Inj.	
706	Venlafaxine	37.5 mg	Tab.	
707	Venlafaxine	75 mg	Tab.	
708	Zuclopenthixol	200 mg	Inj.	1 ml
	<b>RADIOLOGICAL DIAGNOSTICS AGENTS</b>			
709	Barium Sulphate	60% w/v	Liq.	
710	Barium Sulphate	99% w/w	Powder	
711	Dimeglumine Gadopentetate	469 mg/mL	Inj.	
712	Gadodiamide	287mg/0.5mm ol	Inj.	20ml
713	Iohexol	300mgI/ml	Inj.	
714	Iohexol	350mgI/ml	Inj.	
715	Iopamidol	300mgI/ml	Inj.	
716	Iopamidol	370mgI/ml	Inj.	
717	Iopromide	300mgI/ml	Inj.	
718	Iopromide	370mgI/ml	Inj.	
719	Meglumine Iodine	76% w/v 370 mg/ml	Soln.	50 ml

720	Meglumine Iodine	76% w/v 370 mg/ml	Soln.	100 ml
721	Meglumine Iodine	76% w/v 370 mg/ml	Soln.	20 ml
722	Sodium Amidotrizoate (Sodium diatrizoate) + Meglumine Amidotrizoate (Meglumine diatrizoate).	100mg+660mg/ml	Soln.	100ml
<b>RESPIRATORY DRUGS</b>				
723	Acefylline	125 mg /5ml	Syp.	120 ml
724	Aminophylline	25 mg/1ml	Inj.	10 ml
725	Beclomethasone	800 mcg/2ml	Soln.	2 ml
726	Beclomethasone + Salbutamol	50 mcg + 100 mcg	Spray / Inhaler.	
727	Beclomethasone Dipropionate	250 mcg	Inhaler	
728	Budesonide	50 mcg/Actu.	Inhaler	
729	Budesonide	200 mcg	Rota Cap.	
730	Budesonide	400 mcg	Rota Cap.	
731	Budesonide + Formoterol	100 mcg + 6 mcg	Rota Cap.	
732	Budesonide + Formoterol	200 mcg + 6 mcg	Rota Cap.	
733	Budesonide + Formoterol	400 mcg + 6 mcg	Rota Cap.	
734	Diphenhydramine+ Aminophylline+ Ammonium Chloride	8mg+32mg+30 mg /5ml	Syp.	120ml

735	Doxofylline	400mg	Cap.	
736	Doxofylline	100mg/5ml	Syp.	60ml
737	Fluticasone Propionate + Salmeterol	125 mcg + 25mcg	Inhaler	
738	Ipratropium Bromide	20 mcg	Inhaler	
739	Ipratropium Bromide	250 mcg/ml	Soln.	2ml
740	Ketotifen	1 mg	Tab.	
741	Ketotifen	0.2 mg/ml	Syp.	60ml
742	Salbutamol	2 mg	Tab.	
743	Salbutamol	4 mg	Tab.	
744	Salbutamol	2mg/5ml	Syp.	60ml
745	Salbutamol	5mg/ml	Soln.	20ml
746	Salbutamol	100 mcg	Inhaler	
747	Salbutamol	0.5 mg/ml	Inj.	1ml
748	Terbutaline Sulphate	2.5 mg	Tab.	
749	Terbutaline Sulphate	0.3 mg/ml	Syp.	60ml
750	Terbutaline Sulphate	0.5 mg/ml	Inj.	1ml
751	Tiotropium	18 mcg	Rota Cap.	

	<b>STERILE OPHTHALMIC PREPARATIONS</b>			
752	Acyclovir	3% w/w	Eye Oint.	4.5 gm
753	Artificial Tears (Hypromellose + Dextran)	0.3% w/v + 0.1% w/v	Eye Drops	15 ml
754	Acetylcholine	20 mg/ Vial	Inj.	
755	Betamethasone	0.1% w/v	Eye Drops	7.5 ml
756	Brinzolamide + Brimonidine	10mg + 2mg /ml	Eye Drops	5ml
757	Chloramphenicol	1% w/w	Eye Ointment	5gm
758	Chloramphenicol	0.5 % w/v	Eye Drops	10ml
759	Ciprofloxacin	0.3% w/v	Eye Drops	5ml
760	Cyclopentolate	1%	Eye Drops	10ml
761	Cyclopentolate + Proparacaine	1% + 0.5%	Eye Drops	
762	Dexamethasone	0.1% w/v	Eye Drops	
763	Diclofenac Sodium	0.1% w/v	Eye Drops	
764	Dorzolamide + Timolol	2 + 0.5%	Eye Drops	5ml
765	Fluorescein	2% w/v	Eye Drops	15ml
766	Fluorescein	0.6 mg	Strips	
767	Fluorometholone + Neomycin	0.1%+0.5%	Eye Drops	5ml

768	Homatropine	2% w/v	Eye Drops	15ml
769	Latanoprost	0.05%	Eye Drops	2.5ml
770	Levobunolol	0.5% w/v	Eye Drops	5ml
771	Moxifloxacin	0.5% w/v	Eye Drops	5ml
772	Phenylephrine	10 % w/v	Eye Drops	5 ml
773	Pilocarpine HCl	2% w/v	Eye Drops	10 ml
774	Pilocarpine HCl	4% w/v	Eye Drops	10 ml
775	Polymyxin B+ Neomycin + Dexamethasone		Eye Drops	5 ml
776	Polymyxin B+ Neomycin + Dexamethasone		Oint.	3.5 gm
777	Polymyxin B Sulphate + Bacitracin	10,000 IU/gm + 500 IU/gm	Eye Oint.	6 gm
778	Proparacaine	0.5% w/v	Eye Drops	15 ml
779	Ranibizumab	10 mg/ ml	Inj.	
780	Tetracycline	1%	Eye Oint.	5gm
781	Timolol Maleate	0.25%	Eye Drops	5 ml
782	Timolol Maleate	0.5% w/v	Eye Drops	5 ml
783	Tobramycin	0.3% w/v	Eye Drops	5 ml
784	Tobramycin + Dexamethasone	0.3% + 0.1% w/v	Eye Drops	5 ml

785	Travoprost	40mcg/ml	Eye Drops	2.5ml
786	Tropicamide	1% w/v	Eye Drops	15ml
	<b>TOPICAL DRUGS PREPARATIONS</b>			
787	Acyclovir Ointment	5% w/w	Oint.	5 gm
788	Betamethasone dipropionate	0.05%	Oint.	10 gm
789	Betamethasone dipropionate	0.05%	Cream	10 gm
790	Betamethasone dipropionate	0.05%	Lot.	20 ml
791	Benzyl Benzoate	25%	Lot.	120 ml
792	Betamethasone Dipropionate + Gentamicin sulphate	0.05 % + 0.1%	Cream	15 gm
793	Betamethasone Dipropionate + Gentamicin sulphate	0.05 % +0.1 %	Oint.	15gm
794	Calamine	15%	Lot.	120 ml
795	Clobetasol Propionate	0.05% w/w	Cream	10gm
796	Clotrimazole	1%	Cream	10gm
797	Clotrimazole	1%	Lot.	60ml
798	Clotrimazole	1%	Soln.	20ml
799	Coal Tar	4%	Soln.	
800	Fluocinolone Acetonide	0.03%	Cream	15gm

801	Fluocinolone Acetonide	0.03%	Gel	15gm
802	Fusidic acid	2%	Cream	15gm
803	Fusidic acid	2%	Oint.	15gm
804	Gentamicin	0.10%	Cream	10gm
805	Gentamicin	0.10%	Oint.	10gm
806	Gentian Violet	0.50%	Aq. Soln.	
807	Hydrocolloid		Gel	
808	Hydrocortisone	1%	Oint.	10 gm
809	Hydrocortisone	1%	Cream	10 gm
810	Isotretinoin + Erythromycin	0.05 % + 2% w/w	Gel	
811	Lignocaine HCl (Sterile)	2%	Gel	
812	Meglumine antimoniate		Inj.	
813	Miltefosine	10 mg	Tab. / Cap.	
814	Miltefosine	50 mg	Tab. / Cap.	
815	Mupirocin	2 % w/w	Cream	15 gm
816	Mupirocin	2 % w/w	Oint.	15 gm
817	Permethrin	5% w/w	Cream	30gm

818	Permethrin		Lot.	60ml
819	Polymyxin B Sulphate + Bacitracin zinc	10000 IU/g + 500 IU/g	Oint.	10 gm
820	Polymyxin B Sulphate + Bacitracin zinc	10000 IU/g + 500 IU/g	Oint.	20 gm
821	Salicylic Acid	5%	Soln.	
822	Silicone		Gel	
823	Silver Sulfadiazine	1%	Cream	50 gm
824	Silver Sulfadiazine	1%	Cream	250 gm
825	Sodium Stibogluconate		Inj.	
826	Terbinafine	1%	Cream	10gm
827	Terbinafine		Lot.	
828	Tetrachlorodecaoxide	0.052 mg/ 5ml	Soln.	50ml
	<b>DISINFECTANT &amp; ANTISEPTIC</b>			
829	Chloroxylonol	4.80%	Soln.	Various pack sizes one liter and higher volume
830	Chlorhexidine Digluconate	7.10%	Soln.	
831	Formalin Pure	47%	Soln.	450 ml
832	Glutaraldehyde Solution for Sterilization	2%-2.5%	Soln.	5 Litres

833	Hand sanitizer Iso-Propyl Alcohol Based (As per WHO Recommendations)	75%	Soln.	1000ml
834	Hand sanitizer Ethyl Alcohol Based (As per WHO Recommendations)	80%	Soln.	1000ml
835	Hydrogen Peroxide	6%	Soln.	
836	Povidone Iodine	10%	Soln.	450 ml
837	Povidone Iodine	7.5% w/w	Scrub	450 ml
838	Sodium Hypochlorite	10%	Soln.	500 ml
	<b>VITAMINS / MINERALS</b>			
839	Alfacalcidol	0.5 mcg	Tab.	
840	Ascorbic Acid	500 mg	Tab.	
841	Calcium Acetate		Inf.	
842	Calcium Acetate	667mg	Tab.	
843	Calcium Carbonate	(at least containing but not limited to) 327mg	Tab.	
844	Calcium Phosphate	210 mg/ 5ml	Syp.	
845	Cholecalciferol (Vitamin D3)	200000 IU	IM/ Oral Inj.	1ml
846	Pyridoxine HCl	50 mg	Tab.	

847	Retinol (Vitamin A)		Cap.	
	<b>COTTON, BANDAGES, P.O.P, SURGICAL DISPOSABLES &amp; NON-DRUG ITEMS</b>			
848	Absorbable Hemostatic Gelatin Sponges	Different Sizes		
849	Abrams Pleural Biopsy Needles	All sizes		
850	Adhesive Tapes (Paper)	1" width and various lengths		
851	Adhesive Tapes (Paper)	2" width and various lengths		
852	Adhesive Tapes (Paper)	3" width and various lengths		
853	Adhesive Tapes (Paper)	4" width and various lengths		
854	Adhesive Tapes (Plastic)	1" width and various lengths		
855	Adhesive Tapes (Plastic)	2" width and various lengths		
856	Adhesive Tapes (Plastic)	3" width and various lengths		
857	Adhesive Tapes (Plastic)	4" width and various lengths		
858	Angiography Guide Wires	All Sizes		
859	Angiography Exchange Guide Wires	All Sizes		
860	Arterial Catheter (Sterile, wings having holes, Spring-Wire Guide Handle, Black Feed Tube Marker, Return window) The Cannula should be radio- opaque, as well as latex, pyrogen and PVC free)	Different Sizes		

861	Arterial Sheath (Femoral)	All sizes		
862	Automated External Defibrillator			
863	Bacterial Binding Dressing	Different Sizes		
864	Bacterial filter, HME Filter and Viral filter (HCV, HBS+HIV etc.)			
865	Bain Circuit	Adult		
866	Bain Circuit	Pediatric		
867	Bare Metal Cardiac Stents (Cobalt Chromium)	All Sizes		
868	Bare Metal Cardiac Stents (Platinum Chromium)	All Sizes		
869	Bare Metal Cardiac Stents (Stainless Steel)	All Sizes		
870	Becker Implant			
871	Blood Bags (CPDA-1) + Transfusion Sets	Single	500ml	
872	Blood Bags (CPDA-1) + Transfusion Sets	Single	250ml	
873	Blood Bags (CPDA-1) + Transfusion Sets	Double	500ml	
874	Blood Bags (CPDA-1) + Transfusion Sets	Double	250ml	
875	Blood Bags (CPDA-1) + Transfusion Sets	Triple	500ml	

876	Blood Bags (CPDA-1) + Transfusion Sets	Triple	250ml	
877	Blood Collection Tubes (Purple Top)	5 ml		
878	Blood Collection Tubes (Red Top)	5 ml		
879	Blood Collection Tubes (Black Top)	5ml		
880	Blood Collection Tubes (Green Top)	5 ml		
881	Blood Collection Tubes (Yellow Top)	5 ml		
882	Blood Collection Tubes (Blue Top)	5 ml		
883	Blood Collection Tubes (Grey Top)	5 ml		
884	Blood Collection Tubes (White Top)	5 ml		
885	Blood Collection Tubes (Orange Top)	5 ml		
886	Calcium Alginate Dressing	7.5cm x12cm		
887	Calcium Alginate Dressing	10 cm x 20cm		
888	Calcium Alginate Dressing	15cm x 25cm		
889	Calcium Alginate Dressing	Rope 2gm		
890	Casting Tape	6"		
891	Casting Tape	4"		
892	Chest Drainage bottle with Tubing			

893	Chest Tube (with trocar)	Different size		
894	Chest Tube (without trocar)	Different size		
895	Circular Stapler			
896	Colostomy bags (Set comprising bag, adhesive ring, clamp) surfit system			
897	Cord Clamp			
898	Compression face mask			
899	Cotton (Surgical) Corded	200 gm	Roll	
900	Cotton (Surgical) Corded	100 gm	Roll	
901	Cotton Bandages (Surgical)	6.5 cm x 4 m		
902	Cotton Bandages (Surgical)	7.5 cm x 4m		
903	Cotton Bandages (Surgical)	10 cm x 4 m		
904	Cotton Bandages (Surgical)	15 cm x 4 m		
905	Couch Roll	60 cm x 80 m		
906	Condom Catheter	All Sizes		
907	CPAP mask (Continuous positive air pressure mask)	Adult		
908	CPAP mask (Continuous positive air pressure mask)	Pediatric		

909	Crepe Bandages BPC	2.5cm x 4 m		
910	Crepe Bandages BPC	5 cm x 4m		
911	Crepe Bandages BPC	7.5 cm x 4.5 m		
912	Crepe Bandages BPC	15 cm x 4.5 m	Roll	
913	Crepe Bandages BPC	10 cmx4.5m	Roll	
914	CVP line (Single Lumen)	Different Sizes		
915	CVP line (Double Lumen)	Different Sizes		
916	CVP line (Triple Lumen)	Different Sizes		
917	CVP line (Quad Lumen)	Different Sizes		
918	Dental Extraction Forceps			
919	Dental Syringe			
920	Dental wire stainless steel			
921	Diagnostic Catheter	All Types and sizes		
922	Dialysis Catheters (Double Lumen)	16 cmx12F		
923	Dialysis Catheters (Double Lumen)	20 cmx12F		
924	Dialysis Catheters Permanent different sizes	Different size		
925	Disposable Endotracheal Tube without Cuff	2.5 mm		

926	Disposable Endotracheal Tube without Cuff	3 mm		
927	Disposable Endotracheal Tube without Cuff	3.5 mm		
928	Disposable Endotracheal Tube without Cuff	4 mm		
929	Disposable Endotracheal Tube without Cuff	5mm		
930	Disposable Endotracheal Tube without Cuff	5.5mm		
931	Disposable Endotracheal Tube without Cuff	6mm		
932	Disposable Endotracheal Tube without Cuff	6.5mm		
933	Disposable Endotracheal Tube without Cuff	7mm		
934	Disposable Endotracheal Tube without Cuff	7.5mm		
935	Disposable Endotracheal Tube without Cuff	8mm		
936	Disposable Endotracheal Tube with Cuff	4 mm		
937	Disposable Endotracheal Tube with Cuff	4.5 mm		
938	Disposable Endotracheal Tube with Cuff	5mm		
939	Disposable Endotracheal Tube with Cuff	5.5mm		
940	Disposable Endotracheal Tube with Cuff	6mm		
941	Disposable Endotracheal Tube with Cuff	6.5mm		
942	Disposable Endotracheal Tube with Cuff	7mm		

943	Disposable Endotracheal Tube with Cuff	7.5mm		
944	Disposable Endotracheal Tube with Cuff	8mm		
945	Disposable Auto Disable Syringe (Blister packing) sterile	1ml		
946	Disposable Auto Disable Syringe (Blister packing) sterile	3 ml		
947	Disposable Auto Disable Syringe (Blister packing) sterile	5 ml		
948	Disposable Insulin Syringe Ordinary sterile	31 G, 1ml		
949	Disposable Syringe Ordinary (Blister packing) sterile	1ml		
950	Disposable Syringe Ordinary (Blister packing) sterile	3ml		
951	Disposable Syringe Ordinary (Blister packing) sterile	5ml		
952	Disposable Syringe Ordinary (Blister packing) sterile	10ml		
953	Disposable Syringe Ordinary (Blister packing) sterile	20ml		
954	Disposable Syringe Ordinary (Blister packing) sterile	50ml		

955	Disposable Syringe Ordinary (Blister packing) sterile	60ml		
956	Disposable Syringe Ordinary with nozzle for feeding (Blister packing) sterile	60ml		
957	Disposable Sterile Nasogastric Tube	4 Fr		
958	Disposable Sterile Nasogastric Tube	5 Fr		
959	Disposable Sterile Nasogastric Tube	6 Fr		
960	Disposable Sterile Nasogastric Tube	8 Fr		
961	Disposable Sterile Nasogastric Tube	10 Fr		
962	Disposable Sterile Nasogastric Tube	12 Fr		
963	Disposable Sterile Nasogastric Tube	14 Fr		
964	Disposable Sterile Nasogastric Tube	16 Fr		
965	Disposable Sterile Nasogastric Tube	18 Fr		
966	Disposable Sterile Nasogastric Tube	20 Fr		
967	Disposable Sterile Spinal Needle	18 G		
968	Disposable Sterile Spinal Needle	19 G		

969	Disposable Sterile Spinal Needle	20 G		
970	Disposable Sterile Spinal Needle	22 G		
971	Disposable Sterile Spinal Needle	23 G		
972	Disposable Sterile Spinal Needle	25 G		
973	Disposable Tongue depressor wooden			
974	Disposable Dignity Sheet having super absorbency			
975	Disposable Gown (As per WHO standards) • EU PPE Regulation 2016/425 and EU MDD directive 93/42/EEC • FDA class I or II medical device, or equivalent • EN 13795 any performance level, or • AAMI PB70 all levels acceptable, or equivalent			

976	Disposable Sterile Surgical Gloves (without powder) (As per WHO standards) • EU MDD directive 93/42/EEC Category III, • EU PPE Regulation 2016/425 Category III, • EN 455, • ANSI/ISEA 105, • ASTM 6319 or equivalent set of standards	5-9 Sizes		
977	Disposable Sterile Catheter Mount			
978	Disposable suction nozzle			
979	Drill bits	1.2,1.3mm, 1.5mm & 1.6 & 2mm		
980	Drug Eluting Balloon			
981	Drug Eluting Cardiac Stent (Everolimus)	All Sizes		
982	Drug Eluting Cardiac Stent (Sirolimus)	All Sizes		
983	Drug Eluting Cardiac Stents (Zotarolimus)	All Sizes		
984	Disposable OT Cap	Different Sizes		
985	Disposable OT Drapes	Different Sizes		
986	Ear Implant	all sizes		
987	E.C.G sticking Electrodes			
988	Edema compression gloves (Full finger)	Different sizes		

989	Edema compression gloves (Open finger)	Different sizes		
990	Electrosurgical/Diathermy/Cautery Pencil			
991	Epidural kit/ Epidural Anesthesia set Radio-opaque	18 G		
992	Epidural kit/ Epidural Anesthesia set Radio-opaque	20 G		
993	Emergency Cross Head Screws	2.3mm		
994	Emergency Cross Head Screws	2.7mm		
995	Export Aspiration Catheter			
996	Extra Thin Hydrocolloid Dressing	15cm x 15cm		
997	Eye Pads sterile	50x75mm		
998	Eye Pads sterile	55x85mm		
999	Eye Pads sterile	70x54mm		
1000	Eye Pads sterile	57x80mm		
1001	Face Shield (As per WHO standards) • EU PPE Regulation 2016/425, • EN 166, • ANSI/ISEA Z87.1, or equivalent set of standards			
1002	Feeding tube with stopper cap	6 Fr		

1003	Feeding tube with stopper cap	8 Fr		
1004	Feeding tube with stopper cap	10 Fr		
1005	Feeding tube with stopper cap	12 Fr		
1006	Feeding tube with stopper cap	14 Fr		
1007	Feeding tube with stopper cap	16 Fr		
1008	Feeding tube with stopper cap	18 Fr		
1009	Feeding tube with stopper cap	20 Fr		
1010	Fenestrated Silicon Dressing Rolls			
1011	Fiberglass Splint	Different Sizes		
1012	Fistula Cannula (Sterile, small holes along the circumference of end portion, Luer-Lock activated anti-reflux valve & safety cap) The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	Different Gauges		
1013	Fissure Bur			
1014	Flatus Tube	Different Sizes		
1015	Gamgee Wound Dressing	Different Sizes		
1016	Gauze Cutting Scissor	100 cmx 20 m		

1017	Gauze Cloth Roll packing	100 cm x 40 cm		
1018	Gauze Cloth Roll packing			
1019	Gigli Saw (Martensitic steel, two T-shaped handles fitted with a hook on the base end to which a saw wire is attached)	All sizes		
1020	Goggles, protective (As per WHO standards) a. EU PPE Regulation 2016/425, b. EN 166, c. ANSI/ISEA Z87.1, or equivalent			
1021	Guiding Catheter	6 Fr		
1022	Guiding Catheter	7 Fr		
1023	Guide wire for JJ stent	0.25 mm		
1024	Guide wire for JJ stent	0.32 mm		
1025	Guide wire for JJ stent	0.35 mm		
1026	Haemodialyzer with tubing	Adult		
1027	Haemodialyzer with tubing	Paeds		
1028	Hydrogel dressing			
1029	Hydrofiber Dressing	10 cm ×10 cm		

1030	Hydrofiber dressing with silver	20 cm ×30 cm		
1031	Hydrofiber dressing with silver	15 cm×15cm		
1032	Hydrocolloid Dressing	Different sizes		
1033	Irrigation Cannula Stainless steel (Angled)	Different Gauges		
1034	Irrigation Cannula Stainless steel (Straight)	Different Gauges		
1035	Iris Retractor made of bright blue polypropylene, having adjustable silicone stopper (Disposable)			
1036	Intra-aortic Balloon Pump			
1037	I/V fluid administration sets (sterile, minimum 150cm length tubing, latex and pyrogen free, blister pack)			
1038	I/V fluid administration sets (sterile, minimum 150cm length tubing with additional “Y” injection port, latex and pyrogen free, blister pack)			

1039	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	14G		
1040	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	16G		
1041	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	18G		
1042	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	20G		

1043	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	22G		
1044	I/V Cannula (Sterile having wings + injection port in sterilized blister packing. The Cannula should be radio-opaque, as well as latex, pyrogen and PVC free)	24G		
1045	IV Flow Regulator			
1046	Intraosseous Sterile Disposable Infusion Needle, should be latex, pyrogen and PVC free)	Different Gauges		
1047	Infusion Chamber disposable sterile	Adult		
1048	Infusion Chamber disposable sterile	<b>Paeds</b>		
1049	Insulated Nerve Block Needle (Sterile)	21G x 4"		
1050	Isopropyl Alcohol 70% Disposable Nonwoven Swabs			
1051	JJ stent	6FR		

1052	JJ stent	4.7FR		
1053	JJ stent	3.5FR		
1054	K (Kirschner) Wire			
1055	Keratome ophthalmic knife 3.2 mm, 45°			
1056	Laryngeal mask	Different size		
1057	Latex examination gloves Non-Sterile (Powder Free)	Small	Pack of 100 gloves	
1058	Latex examination gloves Non-Sterile (Powder Free)	Medium	Pack of 100 gloves	
1059	Latex examination gloves Non-Sterile (Powder Free)	Large	Pack of 100 gloves	
1060	LP Shunt			
1061	2.7mm Mandible Reconstruction plates (Stainless Steel 316L / 316Lvm /) Titanium) with set	Different sizes and holes		
1062	Manual resuscitator / Self-inflating Bag with Mask	Adult		
1063	Manual resuscitator / Self-inflating Bag with Mask	pediatric		
1064	Manual resuscitator / Self-inflating Bag with Mask	Neonatal		

1065	Medical Shoe Cover (Disposable)			
1066	Disposable Face Mask, (Medical mask, good breathability, and clearly identifiable internal and external faces) (As per WHO or alternative equivalent standards)	<b>Adult</b>		
1067	Malleable Retractor	Different Sizes		
1068	Mucus Extractor			
1069	N-95 (Particulate Respirator) mask As per WHO or Alternative Equivalent standards.	Adult		
1070	Nasal Oxygen Cannula	Neonatal		
1071	Nasal Oxygen Cannula	Pediatric		
1072	Nasal Oxygen Cannula	Adult		
1073	Nebulizer mask with chamber and tubing	Paeds		
1074	Nebulizer mask with chamber and tubing	Adult		

1075	Nitrile Examination Gloves (Powder Free, Non-Sterilized) (As per WHO standards) • EU MDD Directive 93/42/EEC Category III, • EU PPE Regulation 2016/425 Category III, EN 455, EN 374, ANSI/ISEA 105, • ASTM D6319, or alternative equivalent set of standards	Small, Medium and Large		
1076	Non-invasive Ventilation Mask	Different Sizes		
1077	Non-Medicated sterilized adhesive post-operative wound dressing	6x7cm		
1078	Non-Medicated sterilized adhesive post-operative wound dressing	9x10cm		
1079	Non-Medicated sterilized adhesive post-operative wound dressing	9x15cm		
1080	Non-Medicated sterilized adhesive post-operative wound dressing	9x20cm		
1081	Non-Medicated sterilized adhesive post-operative wound dressing	9x25cm		
1082	Non-Medicated sterilized adhesive post-operative wound dressing	9x30cm		

1083	Non-woven Fabric Surgical Adhesive Fix Roll	Various sizes		
1084	Non-rebreather mask	Adult		
1085	Non-rebreather mask	Pediatric		
1086	Nanocrystalline silver dressing	Different Sizes		
1087	Nasal Implant	All Sizes		
1088	Ophthalmic Knife 15°			
1089	Ophthalmic Crescent Knife			
1090	Oxygen Mask	Adult		
1091	Oxygen Mask	Pediatric		
1092	Oropharyngeal Airway	Size 0		
1093	Oropharyngeal Airway	Size 1		
1094	Oropharyngeal Airway	Size 2		
1095	Oropharyngeal Airway	Size 3		
1096	Oropharyngeal Airway	Size 4		
1097	Oropharyngeal Airway	Size 5		
1098	Oropharyngeal Airway	Size 6		
1099	Paraffin Tulle dressing with antiseptic	10x10 cm		

1100	Paraffin Tulle dressing with antiseptic	15x10cm		
1101	Paraffin Tulle dressing with antiseptic	15x150cm		
1102	Partial re-breather mask	Adult		
1103	Partial re-breather mask	Pediatric		
1104	PCI Guide Hydrophilic			
1105	PCI Guide Hydrophobic			
1106	Pigtail with needle for chest drainage and ascitic fluid drainage	Size-14 Size-18, Size-24		
1107	POP Bandages	15 cm x 2.7 m		
1108	POP Bandages	10cm x 2.7 m		
1109	PU Adhesive Incise Drape Film	10 cm x 14 cm		
1110	PU Adhesive Incise Drape Film	15 cm x 28 cm		
1111	PU Adhesive Incise Drape Film	30 cm x 28 cm		
1112	PU Adhesive Incise Drape Film	45 cm x 28 cm		
1113	PU Adhesive Incise Drape Film	55 cm x 44 cm		
1114	Reloadable Linear Cutter Stapler	55mm, 60mm, 75mm, 80 mm staple length		
1115	Scalp Vein Set/ Butterfly Needle/ Winged infusion Set	Different Gauge sizes		

1116	Sterilized disposable needles for dental syringe	Different sizes		
1117	Sterile External Fixators with titanium Alloy Pins	Different Sizes, Shape & Design		
1118	Sterile Nelaton Catheter	12 Fr		
1119	Sterile Nelaton Catheter	14 Fr		
1120	Sterile Nelaton Catheter	16 Fr		
1121	Sterile Skin graft blade for Dermatome Knife	Different Sizes		
1122	Sofra Tulle or at least equivalent			
1123	Spinal Fixation System Full Instrument Set			
1124	Spinal Fusion cage along with pedicle screws and rods	Different sizes		
1125	Silicone rod or Hunter tendon implant	3,4 & 5 mm		
1126	Surgical Saw Stainless steel	All sizes		
1127	Surgical Implants sheets			
1128	Surgical Implants blocks			
1129	Skin Staple Remover			
1130	Skin Stapler Straight			
1131	Steinmann Pins	All Types		

1132	Sterile Gauze Dressing Pad (Radiopaque)			
1133	Sterile Gauze Dressing Pad	10x10 cm		
1134	Sterile Gauze Dressing Pad	15x15 cm		
1135	Sterile Manual Aspirator			
1136	Sterile Suction Catheter	5 Fr		
1137	Sterile Suction Catheter	6 Fr		
1138	Sterile Suction Catheter	8 Fr		
1139	Sterile Suction Catheter	10 Fr		
1140	Sterile Suction Catheter	12 Fr		
1141	Sterile Suction Catheter	14 Fr		
1142	Sterile Suction Catheter	16 Fr		
1143	Sterile Suction Catheter	18 Fr		
1144	Stomahesive Paste			
1145	Stop Cock 3 way with Extension			
1146	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	11		
1147	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	15		

1148	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	22		
1149	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	23		
1150	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	24		
1151	Surgical Blade (Steel carbon, black/ blue/ Stainless Steel)	25		
1152	Suprapubic Catheter			
1153	Thermometer			
1154	Three-Way Foley Catheter	6 Fr		
1155	Three-Way Foley Catheter	8 Fr		
1156	Three-Way Foley Catheter	10 Fr		
1157	Three-Way Foley Catheter	12 Fr		
1158	Three-Way Foley Catheter	14 Fr		
1159	Three-Way Foley Catheter	16 Fr		
1160	Three-Way Foley Catheter	18 Fr		
1161	Three-Way Foley Catheter	20 Fr		
1162	Three-Way Foley Catheter	22 Fr		

1163	Two-Way Foley Catheter 100% Silicon)	6Fr		
1164	Two-Way Foley Catheter 100% Silicon)	8Fr		
1165	Two-Way Foley Catheter 100% Silicon)	10Fr		
1166	Two-Way Foley Catheter 100% Silicon)	12Fr		
1167	Two-Way Foley Catheter 100% Silicon)	14Fr		
1168	Two-Way Foley Catheter 100% Silicon)	16Fr		
1169	Two-Way Foley Catheter 100% Silicon)	18Fr		
1170	Two-Way Foley Catheter 100% Silicon)	20Fr		
1171	Two-Way Foley Catheter 100% Silicon)	22Fr		
1172	Two-Way Foley Catheter (Silicon Coated)	6Fr		
1173	Two-Way Foley Catheter (Silicon Coated)	8Fr		
1174	Two-Way Foley Catheter (Silicon Coated)	10Fr		
1175	Two-Way Foley Catheter (Silicon Coated)	12Fr		
1176	Two-Way Foley Catheter (Silicon Coated)	14Fr		
1177	Two-Way Foley Catheter (Silicon Coated)	16Fr		
1178	Two-Way Foley Catheter (Silicon Coated)	18Fr		
1179	Two-Way Foley Catheter (Silicon Coated)	20Fr		

1180	Two-Way Foley Catheter (Silicon Coated)	22Fr		
1181	Tissue Expander	all types & sizes		
1182	Titanium Micro screw	all sizes		
1183	Titanium microplate with set	1.6mm & 16 holes		
1184	Titanium Mesh	12×6 cm× 0.3mm		
1185	Titanium Mesh	12×6 cm× 1.6mm		
1186	Titanium Mesh	12×6 cm×0.6mm		
1187	Titanium mini plates	2.0mm× 20holes		
1188	Titanium surgical screws	1.6 mm× 5 mm		
1189	Titanium surgical screws	1.6 mm× 6 mm		
1190	Titanium surgical screws	2.0 mm × 7mm		
1191	Titanium surgical screws	2.0× 5.5 to 15mm		
1192	Tracheostomy mask			
1193	Tracheostomy Tube with cuff	Different Sizes		
1194	Tracheostomy Tube without cuff	Different Sizes		
1195	Titanium Ligation Clips	LT 300		
1196	Titanium Ligation Clips	LT 400		

1197	Transparent IV Dressing	Different Sizes		
1198	Tru-cut disposable Biopsy Needles with gun (for solid organs)	Different sizes		
1199	Tyvek Suit (As per WHO or alternative equivalent standards)			
1200	Urine bag with let	2000 ml		
1201	Umbilical Venous Catheter (Sterile)	Different sizes		
1202	Vacuum drainage bottle (closed seal) with tube (Disposable)			
1203	Ventilator Circuit			
1204	Venturi Oxygen Mask with different oxygen concentration venturi valve			
1205	VP Shunt			
1206	Wrist Spanning Plate (screw diameter of 2.5 mm) with set			
1207	Wrist Spanning Plate (2.3 mm locking variable angle screws) with set			
1208	X-Ray film	8x10		

1209	X-Ray film	12x15		
1210	X-Ray film	10x12		
1211	X-Ray film	14x17		
1212	X-ray film CR for closed system of various brand	Different Sizes		
1213	X-ray film CT scan	Different sizes		
1214	X-ray film Dental	Different sizes		
1215	X-ray film for MRI	Different sizes		
1216	X-ray Developer + X-ray Fixer Set			
1217	Zinc oxide adhesive Plaster (Cloth Tape)	2.5 cm x 5m		
1218	Zinc oxide adhesive Plaster (Cloth Tape)	5 cm x 5m		
1219	Zinc oxide adhesive Plaster (Cloth Tape)	7.5 cm x 5m		
1220	Zinc oxide adhesive Plaster (Cloth Tape)	10 cm x 5 m		
	<b>LIST OF SURGICAL SUTURES</b>			
	<b>CATGUT CHROMIC</b>			
	<b>Sutures</b>	<b>Sizes</b>		
1221	20mm, ½ circle round bodied taper point needle, strand length 75cm	4/0		
1222	20mm, ½ circle round bodied taper point needle, strand length 75cm	3/0		

1223	26mm, ½ circle round bodied taper point needle, strand length 75cm	2/0		
1224	30mm, ½ circle round bodied taper point needle, strand length 75cm	2/0		
1225	30mm, ½ circle round bodied taper point needle, strand length 75cm	0		
1226	40mm, ½ circle round bodied taper point needle, strand length 75cm	0		
1227	30mm, ½ circle round bodied taper point needle, strand length 75cm	1		
1228	40mm, ½ circle round bodied taper point needle, strand length 75cm	2		
1229	40mm, ½ circle round bodied taper point needle, strand length 75cm	1		
	<b>BLACK BRAIDED SILK</b>			
	<b>Sutures</b>	<b>USP Sizes</b>		
1230	16mm, ½ circle round bodied taper point needle, strand length 75cm	4/0		
1231	16mm, 3/8 circle reverse cutting needle, strand length 75cm	4/0		
1232	17mm, ½ circle round bodied taper point needle, strand length 75cm	4/0		
1233	24mm, 3/8 circle reverse cutting needle, strand length 45cm	4/0		
1234	30mm, ½ circle round bodied taper point needle, Strand length 75cm	3/0		
1235	26mm, 3/8 circle conventional cutting needle, Strand length 45 cm	3/0		

1236	16mm, ½ circle rounded bodied taper point needle, Strand length 75cm	3/0		
1237	26mm, 3/8 circle reverse cutting needle, Strand length 75cm	2/0		
1238	30mm, 3/8 circle reverse cutting needle, Strand length 75cm	2/0		
1239	30mm, 3/8 circle reverse cutting needle, Strand length 75cm	0		
1240	30mm, ½ circle round bodied needle, Strand length 75cm	0		
1241	26mm, ½ circle reverse cutting needle, Strand length 75cm	0		
1242	36mm, ½ circle reverse cutting needle, Strand length 75cm	1		
1243	30mm, ½ circle round bodied taper point needle, Strand length 75cm	1		
1244	30mm, 3/8 circle reverse cutting needle, Strand length 75cm	1		
1245	40mm, ½ circle round bodied taper point needle, Strand length 75cm	1		
1246	36mm, ½ circle reverse cutting needle, strand length 75cm	1		
1247	40mm, ½ circle round bodied taper point needle, strand length 75cm	2		
1248	45mm, 3/8 circle reverse cutting needle, Strand length 75cm	2		
	<b>POLYGLACTINE 910</b>			
	<b>Sutures</b>	<b>Size</b>		

1249	12mm, 3/8 circle reverse cutting needle, Strand length 45 cm	6/0		
1250	12mm, 3/8 circle reverse cutting needle, Strand length 45 cm	5/0		
1251	22mm, 1/2 circle round bodied taper point, Strand length 70 cm	4/0		
1252	19mm, 3/8 circle reverse cutting needle, Strand length 75cm	4/0		
1253	26mm, 1/2 circle round body taper point, Strand length 70 cm	3/0		
1254	19mm, 3/8 circle reverse cutting needle, Strand length 75cm	3/0		
	<b>POLYGLYCOLIC ACID</b>			
	<b>Sutures</b>	<b>Size</b>		
1255	6mm, 3/8 circle micro-point spatulated double armed needle	8/0		
1256	6mm, 1/4 circle micro-point spatulated double armed needle	8/0		
1257	6mm, 3/8 circle micro-point spatulated double armed needle	6/0		
1258	6mm, 1/4 circle micro-point spatulated double armed needle	6/0		
1259	8mm, 1/4 circle Spatulated double armed needle, Strand length 30cm	6/0		
1260	17mm, 1/2 circle round bodied taper point needle, Strand length 75cm	5/0		

1261	16mm, 3/8 circle reverse cutting needle, Strand length 75cm	4/0		
1262	22mm, ½ circle round bodied taper point needle, Strand length 75cm	4/0		
1263	17mm, ½ circle round bodied needle, Strand length 75cm	4/0		
1264	16mm, 3/8 circle reverse cutting needle, Strand length 75 cm	3/0		
1265	22mm, ½ circle round bodied taper point 75 cm	3/0		
1266	26mm, ½ circle reverse cutting needle, Strand length 75 cm	2/0		
1267	30mm, ½ circle round bodied taper point needle, strand length 75 cm	2/0		
1268	36mm, ½ circle round bodied taper point needle, Strand length 90 cm	2/0		
1269	30mm, ½ circle round bodied taper point needle, strand length 75 cm	0		
1270	40mm, ½ circle round bodied taper point needle, strand length 90 cm	0		
1271	30mm, ½ circle round bodied taper point needle, strand length 75cm	1		
1272	40mm, ½ circle round bodied taper point needle, strand length 75 cm	1		
1273	36mm, ½ circle round bodied taper cutting needle, strand length 90 cm	1		

1274	48mm, ½ circle round bodied taper point needle, Strand length 90 cm	2		
1275	40mm, ½ circle round bodied taper point needle, strand length 90cm	2		
	<b>POLYPROPYLENE</b>			
	<b>Sutures</b>	Size		
1276	8mm, 1/2 circle round bodied taper point double armed needle, strand length 60cm	12/0		
1277	8mm, 3/8 circle round bodied taper point double armed needle, strand length 60 cm	11/0		
1278	8mm, 1/2 circle round bodied taper point needle, strand length 60cm	10/0		
1279	12mm, 3/8 circle reverse cutting needle, strand length 45 cm	9/0		
1280	16mm, 3/8 circle curved cutting, strand length 90 cm	8/0		
1281	13mm, 1/2 circle round bodied taper point double armed needle, strand length 75cm	6/0		
1282	12mm, 3/8 circle reverse cutting needle, strand length 45cm	6/0		

1283	16mm, 3/8 circle curved cutting needle, strand length 90cm	6/0		
1284	10mm, 1/2 circle round bodied taper point double armed needle, strand length 60cm	6/0		
1285	13mm, 1/2 circle round bodied taper point double armed needle, strand length 75cm	5/0		
1286	16mm, 1/2 circle round bodied taper point double armed needle, strand length 90cm	5/0		
1287	13mm, 1/2 circle curved cutting needle, strand length 90cm	5/0		
1288	16mm, 1/2 circle curved cutting, strand length 90cm	5/0		
1289	15mm, 1/2 circle round bodied taper point needle, strand length 75cm	4/0		
1290	16mm, 1/2 circle round bodied taper point double armed, strand length 90 cm	4/0		
1291	19mm, 3/8 circle reverse cutting needle, strand length 75cm	4/0		

1292	16mm, 1/2 circle round bodied taper point double armed needle, strand length 90cm	3/0		
1293	20mm, 1/2 circle curved cutting, strand length 90cm	3/0		
1294	26mm, 1/2 circle curved cutting needle, strand length 90cm	3/0		
1295	24mm, 3/8 circle reverse cutting needle, strand length 45cm	3/0		
1296	16mm, 1/2 curved cutting needle, strand length 90cm	3/0		
1297	25mm, 1/2 circle round bodied taper point double armed needle, strand length 90cm	3/0		
1298	30 mm, 1/2 circle round bodied taper point double armed needle, strand length 90cm	3/0		
1299	30mm, 1/2 circle round bodied taper point, strand length 75cm	2/0		
1300	75mm, straight cutting needle, 75cm	2/0		

1301	25mm, 1/2 circle round bodied taper point, strand length 90cm	2/0		
1302	25mm, 1/2 circle round bodied taper cut point needle, strand length 75 cm	2/0		
1303	36mm, 1/2 circle reverse cutting needle, strand length 75cm	0		
1304	30mm, 1/2 circle round bodied taper point needle, strand length 75cm	0		
1305	30mm, 1/2 circle round bodied taper point needle, strand length 75cm	1		
	<b>POLYPROPYLENE MESH</b>			
	<b>Mesh</b>	<b>Sizes</b>		
1306	Polypropylene Mesh	30cm x 30cm		
1307	Polypropylene Mesh	15 cm x 15cm		
1308	Polypropylene Mesh	15 cm x 6cm		
1309	Polypropylene Mesh	6cm x 11cm		
	<b>POLYAMIDE</b>			
	<b>Suture</b>	<b>Sizes</b>		
1310	6.5 mm, 3/8 circle spatula micro point needle, strand length 30cm	10/0		

	<b>POLYESTER</b>			
	<b>Sutures</b>	Sizes		
1311	24mm, ½ circle round bodied needle, strand length 75cm	2/0		
	<b>POLYDIOXANONE</b>			
1312	13mm, 3/8 circle round bodied taper point double armed needle, strand length 45cm	7/0		
1313	13mm, ½ circle round bodied taper point needle, strand length 75cm	6/0		
1314	17mm, ½ circle round bodied taper point needle, strand length 75cm	6/0		
1315	13mm, 3/8 circle round bodied taper point double armed needle, strand length 75cm	6/0		
1316	13mm, ½ circle round bodied taper point needle, strand length 75cm	5/0		
1317	17mm, ½ circle round bodied taper point needle, strand length 75cm	5/0		
1318	13mm ½ circle round bodied taper point double armed needle, strand length 75cm	5/0		

1319	17mm, 3/8 circle round bodied taper point double armed needle, strand length 75cm	5/0		
1320	19mm 3/8 circle round bodied taper point double armed needle, strand length 75cm	5/0		
1321	12mm, 3/8 circle reverse cutting needle, strand length 75cm	5/0		
1322	19mm, 3/8 circle reverse cutting needle, strand length 75cm	5/0		
1323	17mm, 1/2 circle round bodied taper point needle, strand length 75cm	4/0		
1324	22mm, 1/2 circle round bodied taper point needle, strand length 75cm	4/0		
1325	26mm, 1/2 circle round bodied taper point needle, strand length 75cm	4/0		
1326	17mm, 1/2 circle round bodied taper point double armed, strand length 75cm	4/0		
1327	17mm, 1/2 circle round bodied	3/0		

1328	taper point needle, strand length 75cm			
1329	22mm, 1/2 circle round bodied taper point needle, strand length 75cm	3/0		
1330	26mm, 1/2 circle round bodied taper point needle, strand length 75cm	3/0		
1331	30mm, 1/2 circle round bodied taper point needle, strand length 75cm	3/0		
1332	19mm, 3/8 circle reverse cutting needle, strand length 75cm	3/0		
1333	26mm 3/8 circle reverse cutting needle, strand length 75cm	3/0		
1334	26mm, 1/2 circle round bodied taper point needle, strand length 75cm	2/0		
1335	30mm, 1/2 circle round bodied taper point needle, strand length 75cm	2/0		
1336	36mm, 1/2 circle round bodied taper point needle, strand length 75cm	2/0		

1337	40mm, 1/2 circle round bodied taper point needle, strand length 75cm	2/0		
1338	26mm, 3/8 circle reverse cutting needle, strand length 75cm	2/0		
1339	30mm, 3/8 circle reverse cutting needle, strand length 75cm	2/0		
1340	36mm 1/2 circle reverse cutting needle, strand length 75cm	2/0		
1341	36mm, 1/2 circle round bodied taper point needle, strand length 75cm	1		
1342	40mm, 1/2 circle round bodied taper point needle, strand length 75cm	1		
1343	48mm, 1/2 circle round bodied taper point needle, strand length 75cm	1		
1344	40mm, 3/8 circle reverse cutting needle, strand length 90cm	1		
1345	36mm, 1/2 circle reverse cutting needle, strand length 75cm	1		

1346	26mm, 1/2 circle round bodied taper point needle	1		
1347	30mm, 1/2 circle round bodied taper point needle	0		
1348	36mm, 1/2 circle round bodied taper point needle	0		
1349	40mm, 1/2 circle round bodied taper point needle	0		
1350	24mm 3/8 circle reverse cutting needle, strand length 75cm	0		
1351	36mm, 1/2 circle reverse cutting needle, strand length 75cm]	0		
	<b>NYLON SUTURES</b>			
	<b>Sutures</b>	<b>Size</b>		
1352	6 mm, 3/8 circle micro point spatula double needle, strand length 30cm	10/0		
1353	6.2mm, 3/8 circle micro point spatula double needle, strand length 30cm	10/0		
	<b>STAINLESS STEEL SUTURES/ WIRE</b>			

	<b>Sutures</b>	<b>Sizes</b>		
1354	48mm, ½ circle round bodied tapercut point needle, strand length 45cm	<b>5</b>		
1355	48mm, ½ circle round bodied tapercut point needle, strand length 45cm	<b>4</b>		
	<b>BONE WAX, CEMENT &amp; GRANULES</b>			
1356	Antibiotic-impregnated bone cement			
1357	Bone Substitute Granules	0.5cc &10cc		
1358	Bone Wax			
1359	Bone cement			

## List of Abbreviation

S.N	Words	Abbreviations
1.	Actuation	Actu.
2.	Aqueous	Aq.
3.	Capsule	Cap.
4.	Cartridges	Ctg.
5.	Centimeter	Cm
6.	Citrate Phosphate Dextrose Adenine-1	CPDA-1
7.	Dispersible	Disper.
8.	Enteric Coated	EC.
9.	Extended-release Tablet	ER-Tab.
10	French Gauge	F / Fr
11	Gram	gm
12	Gauge	G
13	Infusion	Inf.
14	Inhalation	Inh.
15	Injection	Inj.
16	Intramuscular	IM
17	Intravenous	IV
18	International Unit	IU
19	Liquid	Liq.
20	Liter	L
21	Lotion	Lot.
22	Meter	m
23	Microgram	mcg
24	Milligram	mg

25	Milliliter	ml
26	Millimeter	mm
27	Million International Unit	MIU
28	Millimole	mmol
29	Ointment	Oint.
30	Operation theatre Cap	OT Cap
31	Operation theatre Drape	OT Drape
32	Quadruple	Quad.
33	Solution	Soln.
34	Sublingual Tablet	SL. Tab.
35	Suppository	Supp.
36	Suspension	Susp.
37	Sustained Release	SR-Tab.
38	Syrup	Syp.
39	Tablet	Tab.
40	United States Pharmacopeia	USP
41	Vaginal Tablet	Vag. Tab.
42	Weight/ Weight	w/w
43	Weight/Volume	w/v

## **Section V. Technical Specifications**

### **Technical Evaluation Criteria for Drugs / Medicines, Medical Devices, Surgical Disposables and Non-Drug Items (NDIs)**

**(Maximum Allocable Marks Score for Technical Evaluation = 70 Marks)**

**NOTE:**

*For further details of evaluation criteria and marking scheme, please see relevant proformas for technical evaluation of these SBDs.*

**1. SYSTEM BREAKING / DISQUALIFICATION POINTS IN TECHNICAL EVALUATION CRITERIA:**

- a. These system breaking / disqualification points mentioned in this section are in addition to the provision of mandatory documents, as elaborated in Bid Cover Sheet (Bid Form-1).
- b. During technical evaluation of the quoted bids, bidders may stand disqualified if the Scrutiny Committee for bids evaluation and /or Inspection Team/s find and declare any of the shortcoming/s related to the documents and/or manufacturing units and / or the premises of the manufacturers and /or Importers regardless of completion / fulfillment or otherwise of any terms and conditions, criteria and /or codal formalities.
- c. The technical & financial evaluation system for Govt: MCC bids for the FY 2021-23 comprises fifteen different evaluation proformas (Section V. Technical Specifications) each having system breaking points and non-compliance of any of these system breaking parameters on part of bidder shall lead to disqualification of firm and /or quoted item/s, whatever the case may be.
- d. Further details of system breaking points / issues for various categories of items are as follows:
  - A. **Manufacturer of General Drugs/Medicines, I/V Fluids and Powdered Injectable Drugs:**
    - i. **Stability chamber:**  
Non-availability and / or Non-functioning of Stability Chambers (Both Accelerated and Real time) due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm.
    - ii. **Raw Material Store / Storage:**  
Nonadherence to cGMP and / or cGSP due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
    - iii. **Finished Goods Store / Storage:**  
Nonadherence to GSP, due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
    - iv. **Functional HVAC:**  
Non-availability or non-functioning of HVAC system in relevant areas of the factory due to any reason, whatsoever, at the time of inspection

by the MCC expert/s shall lead to disqualification of the firm from this bidding competition. Non-Availability of adequate, qualified and relevant Human Resource due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

**B. Importers of General Drugs/Medicines, I/V Fluids and Powdered Injectable Drugs:**

- i. Valid cGMP/Certificate of Pharmaceutical Product (COPP)/ Certificate of Medicinal Product (COMP)/ Certificate of the Principal Manufacturer for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- ii. Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20% stock at the time of inspection shall lead to disqualification of the quoted item/s).
- iii. Functional and effective Air-conditioning & Ventilation System and effective cold chain (thermo-labile drugs). Adherence to Good storage practices (GSP). Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.
- iv. Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- v. Valid cGMP (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin) in **original** and Valid Free sale certificate for the quoted item/s duly attested by the Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan in **original** shall be provided to the Inspection team at the time of inspection.

**C. Manufacturer of Biological Products:**

- i. Stability chamber:  
Non-availability and / or Non-functioning of Stability Chambers (Both Accelerated and Real time) due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm.
- ii. Raw Material Store / Storage:  
Non adherence to cGMP and / or cGSP due to any reason, whatsoever,

at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

iii. Finished Goods Store / Storage:

Non adherence to GSP, due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

iv. Functional HVAC:

Non-availability or non-functioning of HVAC system in relevant areas of the factory due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

v. Non-Availability of adequate, qualified and relevant Human Resource due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

vi. Non-Availability of Functional and Effective Cold Chain System & Uninterrupted Power Supply in all areas of raw material storage, finished goods storage, in-process quarantine and production (evaluated by the panel of expert at the time of inspection and non-adherence to cGMP) shall lead to disqualification of the firm.

**D. Importer/s of Biological Products:**

i. Valid cGMP Certificate of the Principal Manufacturer for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.

ii. Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s).

iii. Functional and effective Air-conditioning & Ventilation System and effective cold chain (thermo-labile drugs) non provision of the facility will lead to Disqualification.

iv. Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.

v. Valid cGMP (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin) in **original** and Valid Free sale certificate for the quoted item/s duly attested by the

Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan in **original** shall be provided to the Inspection team at the time of inspection.

**E. Manufacturer/s of Medical Devices (excluding Cardiac Stents):**

- i. Valid cGMP certificate issued by DRAP.
- ii. Raw Material Store / Storage:  
Non adherence to cGMP and / or cGSP due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
- iii. Finished Goods Store / Storage:  
Non adherence to GSP, due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
- iv. Functional HVAC:  
Non-availability or non-functioning of HVAC system in relevant areas of the factory due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
- v. Non-Availability of adequate, qualified and relevant Human Resource due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.
- vi. Quality assurance certificate for the quoted devices from any international accredited body/DRAP.
- vii. Samples of devices will be tested and evaluated by the Drugs Testing Laboratory as well as by a panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.

**F. Importer/s of Medical Devices (excluding Cardiac Stents):**

- i. Valid cGMP /Quality Control /Quality Assurance Certificate of the Principal Manufacturer from an IAF accredited body or any accredited / authorized / regulatory body in the country of origin for the quoted item/s as issued for the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.  
(In case of Non-applicability of the above mentioned certificates for Adhesive Tape (Non Sterile) only, provision of EC-Declaration of conformity from the principal manufacturer (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s is mandatory).
- ii. Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20% stock at the time of inspection shall lead to

- disqualification of the quoted item/s).
- iii. Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.
- iv. Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- v. Samples of devices will be tested and evaluated by the Drugs Testing Laboratory as well as by a panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.
- vi. Valid cGMP /Quality Control /Quality Assurance Certificate (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin) in **original** and Valid Free sale certificate for the quoted item/s duly attested by the Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan in **original** shall be provided to the Inspection team at the time of inspection.

**G. Importer/s of Medical Devices (Cardiac Stents)**

- i. Valid cGMP /Quality Control /Quality Assurance Certificate of the Principal Manufacturer for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- ii. Valid certification of US Food and Drug Administration (US FDA) of quoted item/s. Non-provision of this certificate shall lead to disqualification of the quoted item/s.
- iii. Valid permission of sale or import of quoted item/s for sale in the US open market. Non-provision of this certificate shall lead to disqualification of the quoted item/s.
- iv. Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20% stock at the time of inspection shall lead to disqualification of the quoted item/s)
- v. Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the

firm.

- vi. Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- vii. Samples of devices will be tested and evaluated by the Drugs Testing Laboratory as well as by a panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.
- viii. Valid cGMP /Quality Control /Quality Assurance Certificate (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin) in **original** and Valid Free sale certificate for the quoted item/s duly attested by the Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan in **original** shall be provided to the Inspection team at the time of inspection.

**H. Manufacturer/s of Cotton & Related Goods:**

- i. Functional and effective Air-conditioning & Ventilation System (evaluated by the panel of expert, Non functionality of the Air-conditioning & Ventilation system in specified section/s shall lead to disqualification of the section or firm).
- ii. Appropriate storage of raw material/s as per law (evaluated by the Inspection Team/s). Non provision of good storage condition shall lead to disqualification of the section or firm.
- iii. Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s, shall lead to Disqualification of the firm.
- iv. Samples of devices will be tested and evaluated by the Drugs Testing Laboratory as well as by a panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.
- v. Non-availability of adequate, qualified and relevant Human Resource due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

**I. Manufacturers of Non-Drug Items:**

- i. Raw material storage (as evaluated at the time of inspection by the MCC expert/s). Non adherence to cGMP shall lead to disqualification of the firm.
- ii. Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s, shall lead to Disqualification of the firm.
- iii. Functional HVAC (as evaluated by the MCC expert/s at the time of

inspection). Non functionality of the HVAC system shall lead to Disqualification of the relevant section/firm.

- iv. Samples of devices will be tested and evaluated by the Drugs Testing Laboratory as well as by a panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.
- v. Non-Availability of adequate, qualified and relevant Human Resource due to any reason, whatsoever, at the time of inspection by the MCC expert/s shall lead to disqualification of the firm from this bidding competition.

**J. Importer/s of Non-Drug Items:**

- i. Valid cGMP /Quality Control /Quality Assurance Certificate of the Principal Manufacturer from an IAF accredited body or any accredited / authorized / regulatory body in the country of origin for the quoted item/s as issued for the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.  
(In case of Non-applicability of the above mentioned certificates for Examination Gloves (Non Sterile) only, provision of EC-Declaration of conformity from the principal manufacturer (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s is mandatory).
- ii. Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20% stock at the time of inspection shall lead to disqualification of the quoted item/s).
- iii. Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.
- iv. Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm.
- v. Samples of devices will be tested by the panel of experts / end users and the quoted item/s may be disqualified for further competition on the report/s of these entities.

- vi. Valid cGMP /Quality Control /Quality Assurance Certificate (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin) in **original** and Valid Free sale certificate for the quoted item/s duly attested by the Pakistani embassy in the country of origin of quoted item/s or embassy of the country of origin in Pakistan in **original** shall be provided to the Inspection team at the time of inspection.

## Section V. Technical Specifications (Continued)

### **Financial Evaluation and Scoring System for Bids**

**(Maximum Allocable Marks Score = 30 marks)**

The financial bids of technically qualified bidders will be opened publicly at the time to be announced by the Procuring Agency and the financial bids found technically non-responsive shall be returned un-opened to the respective Bidders.

Total Allocable marks for Technical Proposal = 70

Total Allocable marks in Financial Proposal = 30

Total Combined Allocable Score for individual bids = Marks obtained in Technical Evaluation  
+ Marks obtained in Financial Evaluation = 100

#### **Scoring Methodology:**

Contract will be awarded to the lowest evaluated responsive firm whose product ranks highest in the Combined Evaluation scoring calculated through the Marks awarded to Technical Proposal and Financial Proposal as stated in the Bid Data Sheet of these SBDs.

The Evaluation Methodology is a combination of non-price factors (in Technical Criteria) and price factor (in Financial Criteria); and each having points as elaborated in the evaluation proformas provided in these SBDs.

As evident from allocable score above and because of the importance and complexities/sensitivities in the field of procurement and use of Drugs and other products related to human lives and health, this Methodology puts greater emphasis on non-price factors like high quality of the product derived from excellent-grade raw material, stringent product certifications, international best pharmaceutical quality control practices in laboratories, Pharmaco-vigilance systems for Drug safety reporting and monitoring; and the most efficient industrial processes in the manufacturing premises.

**Procedure for the Marks Scoring:** Marks will be awarded or otherwise for various technical parameters to each quoted product based on the prescribed Technical and Financial criteria. The total combined marks will determine the highest-ranking product in each product category for contract award.

**The formula to calculate the marks for the price by the bidders other than lowest bidder is given below:**

Financial Evaluation Score of individual quoted Product:

= [Lowest quoted Price of the item ÷ Next higher proposed Price of the competing item] X Total allocable financial score

#### **Solved Example of Financial Scoring:**

- If the lowest quoted price of an item is Rs. 86/-, the same lowest bidder will obtain score as below:  
=  $[86 \div 86] \times 30$   
= 30 marks, being the lowest bidder for the quoted item.
- If the next higher quoted price of the same item is Rs. 105/-, the marks obtained will be:  
=  $[86 \div 105] \times 30 = 24.57$  Marks
- If the next higher quoted price of the same item is Rs. 130/-, the marks obtained will be:

$= [86 \div 130] \times 30 = 19.84$  Marks  
.... And so on.

ALL TECHNICAL EVALAUTION PROFORMAS IN MS-EXCEL FORMAT ARE AVAILABLE ON OFFICAL WEBSITE OF HEALTH DEPARTMENT. ([WWW.HEALTHKP.GOV.PK](http://WWW.HEALTHKP.GOV.PK), [WWW.DGHSKP.GOV.PK](http://WWW.DGHSKP.GOV.PK) )

Evaluation Criteria for Manufacturers of General Medicine/Drugs Government MCC 2021-23																											
Name of Firm					Technical Evaluation Matrix																				Total Product Evaluated Score		Total Technical Score
S. No.	Product General Information				Factory Technical Evaluation Parameters										Total Factory Evaluated Score	Product Evaluation Parameters										Total Product Evaluated Score	Total Technical Score
					Documents Based Factory Score					Factory Evaluation Visit Score						Product Technical Parameters											
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25		
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 18001/45001 certification)	Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 14001 certification)	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 9001 certification)	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies or Regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificates for equipment in the factory awarded by any PNAC accredited body (duly attested by the senior executive of the firm).	Functional Stability Chamber (Both Accelerated and Real Time) (evaluated at the time of inspection by the MCC experts, as non-availability or non-functioning of Stability Chamber shall lead to disqualification of the firm).	Raw material storage (as evaluated at the time of inspection by the MCC experts). Non adherence to cGMP shall lead to disqualification of the firm.	Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC experts, shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by the MCC experts at the time of inspection, Non-availability or non-functionality of the HVAC system shall lead to Disqualification of the relevant section/firm.	Available and Functional HVAC (as evaluated by the MCC experts at the time of inspection). Non-availability or non-functionality of the HVAC system shall lead to Disqualification of the relevant section/firm.		Current export certificate of the quoted item/s from DRAP not older than one year (certificate duly attested by senior executive of the firm). (COPP/ or export NOC issued by DRAP shall be considered) Export to non SRA countries will be awarded 1 mark per country, maximum 5 marks). Export to any SRA country shall be awarded 5 marks).	Goods Declaration certificate of imported API of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medc or Health Canada or by regulatory authority of SRA countries coupled with Form 3 (form of undertaking to accompany an application for License to import Drugs) and Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of API from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 17, duly attested by the senior executive of the firm.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Valid Certificate of Analysis of the Type / class of material used for the immediate container of the quoted item/s, as issued by the manufacturer of this material coupled with Invoice/proof of purchase:  For award of marks, the certificate of analysis must clearly mention: 1. Pharma Grade Aluminium Foil, PVC, Capsule Shell, Plastic, HDPE or any other material used for the immediate container. 2. Type of Glass material for Liquid ampoules must be USP class 1. 3. Type of Glass material for Oral Sympo/ Suspensions must be USP Type 3 or better.  (Documents duly attested	Stability studies of quoted item/s not older than 3 years (duly attested by the Q/C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks				
	Ref. No. of Item	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	3	3	5	4	4	3	4	4	34	5	5	5	5	3	3	5	5	36	70		

Evaluation Criteria for Importers of General Medicines/Drugs Government MCC 2021-23																											
Name of Firm																											
S. No.	Product General Information				Principal's and Importer's Evaluation Parameters								Suppliers Technical Score	Product Technical Evaluation												Product Evaluated Score	Total Technical Score
					Principal Manufacturer Evaluation				Importer's Evaluation					Product Technical Parameters													
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24			
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 18001/45001 certification	Valid ISO 14001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 14001 certification.	Valid ISO 9001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 9001 certification	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies regulatory body in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificate for equipment in the factory (duly attested by the senior executive of the firm).	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC experts). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s). The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.	Adherence to Good Storage Practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC experts at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by the MCC experts at the time of inspection).	Valid Proof of export by the Principal Manufacturer to US/Europe/SRA countries/ies, not older than one year (certificate duly attested by senior executive of the firm).  03 marks for export to SRA Countries. Export Certificate/ COPP /COMP issued by the relevant regulatory body of the countries mentioned above.	Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medc or Health Canada or by regulatory authority of SRA's countries coupled with Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Valid certificate of the availability of the quoted item in the US market.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Valid Certificate of Analysis of the Type / class of material used for the immediate container of the quoted item/s, as issued by the manufacturer of this material coupled with Invoice/proof of purchase:  For award of marks, the certificate of analysis must clearly mention: 1. Pharma Grade Aluminium Foil, PVC, Capsule Shell, Plastic, HDPE or any other material used for the immediate container. 2. Type of Glass material for Liquid ampoules must be USP class 1. 3. Type of Glass material for Oral Sympo/ Suspensions must be USP Type 3 or better.	Stability studies of quoted item/s not older than 3 years (duly attested by the Q/C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2 - 5 % market share = 3 marks More than 5 % market share = 5 marks						
	Ref. No. of Item	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	3	3	5	7	5	7	34	3	5	5	5	2	3	4	4	5	36	70			

Evaluation Criteria for Manufacturers of General Medicine/Drugs (I/V Fluids) for Government MCC 2021-23																											
Name of Firm																											
S. No.	Product General Information				Factory Technical Evaluation Parameters										Technical Evaluation Matrix												
					Documents Based Factory Score					Evaluation visit Score					Factory Evaluated Score	Product Evaluation Parameters										Product Evaluated Score	Total Techni cal Score
																Product Technical Parameters											
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 18001/45001 certification)	Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 14001 certification)	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 9001 certification)	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies or Regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificates for equipment in the factory awarded by any PNAC accredited body (duly attested by the senior executive of the firm).	Availability Functional Stability Chamber (Both Accelerated and Real Time) (evaluated at the time of inspection by the MCC experts), as non-availability or non-functioning of Stability Chamber shall lead to disqualification of the firm).	Raw material storage (as evaluated at the time of inspection by the MCC experts). Non adherence to cGMP shall lead to disqualification of the firm.	Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC experts, shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC experts) at the time of inspection, Non-availability shall lead to disqualification of the section/s or firm).	Available and Functional HVAC (as evaluated by the MCC experts at the time of inspection). Non-availability or non-functionality of the HVAC system shall lead to Disqualification of the relevant section/firm.	31	Current export certificate of the quoted item/s from DRAP not older than one year (certificate duly attested by senior executive of the firm). COPP or export NOK issued by DRAP shall be considered) Export to non SRA countries will be awarded (1 mark per country, maximum 5 marks). Export to any SRA country shall be awarded 5 marks).	Goods Declaration certificate of imported API of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medc or Health Canada or by regulatory authority of SRAs countries coupled with Form 3 (form of undertaking to accompany an application for License to import Drugs) and Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of API of the quoted item/s from the Principal Manufacturer as in column 16, duly attested by the senior executive of the firm.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Valid Certificate of Analysis of the Type / class of material used for the immediate container of the quoted item/s, as issued by the manufacturer of this material coupled with Invoice/proof of purchase: For award of marks, the certificate of analysis must clearly mention: 1. Pharma Grade Aluminium Foil, PVC, Capsule Shell, Plastic, HDPE or any other material used for the immediate container. 2. Type of Glass material for Liquid ampoules (infusion bottles must be USP class 1. 3. Type of Glass material for Oral Syringe/ Suspensions must be USP Type 3 or better.	I/V fluid container/bottles with rubber stopper & tear off protective seal.	Stability studies of quoted item/s not older than 3 years (duly attested by the Q.C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks	39	70	
	Ref. No. of item in MCC Formulary	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	2	3	5	4	4	3	2	4	31	5	5	5	5	3	4	3	4	5	39	70	

Evaluation Criteria for Import of General Medicines/Drugs I/V fluids for Government MCC 2021-23																										
Name of Firm					Technical Evaluation Matrix										Product Technical Evaluation										Product Evaluated Score	Total Technical Score
S. No.	Product General Information				Principal's and Importer's Evaluation Parameters								Suppliers Technical Score	Product Technical Parameters										Product Availability		
					Principal Manufacturer Evaluation				Importer's Evaluation					Product Technical Parameters												
					1	2	3	4	5	6	7	8		9	10	11	12	13	14	15	16	17	18			
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 18001/45001 certification	Valid ISO 14001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 14001 certification.	Valid ISO 9001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 9001 certification	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificate for equipment in the factory (duly attested by the senior executive of the firm).	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC experts). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s). <b>The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.</b>	Functional and effective Airconditioning & Ventilation System and effective cold chain (thermostable drugs). Adherence to Good storage practices (GSP). Non adherence to GSP, as evaluated by the MCC experts at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC experts) at the time of inspection.	Valid Proof of export by the Principal Manufacturer to US/Europe/SRA country/ies, not older than one year (certificate duly attested by senior executive of the firm).  10 marks for export to SRA Countries. Export Certificate/ COPP/COMP issued by the relevant regulatory body of the countries mentioned above.	Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medc or Health Canada, or by regulatory authority of SRAs countries coupled with Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Valid Certificate of Analysis of the Type / class of material used for the immediate container of the quoted item/s, as issued by the manufacturer of this material coupled with Invoice/proof of purchase: For award of marks, the certificate of analysis must clearly mention: 1. Pharma Grade Aluminium Foil, PVC, Capsule Shells, Plastic, HDPE or any other material used for the immediate container. 2. Type of Glass material for Liquid ampoules must be USP class 1. 3. Type of Glass material for Oral Syringe/ Suspensions must be USP Type 3 or better.  (Documents duly attested	Valid certificate of the availability of the quoted item in the US market.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	I/V fluid container/bottles with rubber stopper & tear off protective seal.	Stability studies of quoted item/s not older than 3 years (duly attested by the Q/C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks				
	Ref. No. of item	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	2	3	5	6	5	6	31	3	5	5	5	4	2	3	3	4	5	39	70	

Evaluation Criteria for Manufacturers of Dry Powder Injectables, Government MCC 2021-23																										
Name of Firm																										
S. No.	Product General Information				Factory Technical Evaluation Parameters										Technical Evaluation Matrix											
					Documents Based Factory Score					Evaluation visit based Score					Factory Evaluated Score	Product Evaluation Parameters										Product Evaluated Score
	1	2	3	4	5	6	7	8	9	10	11	12	13	14		15	16	17	18	19	20	21	22	23	24	
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured, issued by PNAC/ accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 18001/45001 certification)	Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by PNAC/ accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 14001 certification)	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by PNAC/ accredited body (duly attested by senior executive of the firm)  Provide Online verification link of ISO 9001 certification)	Valid accreditation of manufacturing unit or its relevant sections by the US-FDA or WHO or official accreditation body/ies/ or Regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificates for equipment in the factory awarded by any PNAC/ accredited body (duly attested by the senior executive of the firm).	Functional Stability Chamber (Both Accelerated and Real Time) (evaluated at the time of inspection by the MCC expert/s, as non-availability or non-functioning of Stability Chamber shall lead to disqualification of the firm).	Raw material storage (as evaluated at the time of inspection by the MCC expert/s). Non adherence to GMP shall lead to disqualification of the firm.	Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s, shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection, Non-availability shall lead to disqualification of the section/s or firm).	Available and Functional HVAC (as evaluated by the MCC expert/s at the time of inspection). Non-availability or non-functionality of the HVAC system at the time of evaluation visit shall lead to Disqualification of the relevant section/firm.	82	Current export certificate of the quoted item/s from DRAP not older than one year (certificate duly attested by senior executive of the firm). (COPP/ or export NOC issued by DRAP shall be considered) Export to non SRA countries will be awarded (1 mark per country, maximum 5 marks). Export to any SRA country shall be awarded 5 marks).	Goods Declaration certificate of imported API of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by regulatory authority of SRAs countries coupled with Form 3 (form of undertaking to accompany an application for License to import Drugs) and Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of API from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 17, duly attested by the senior executive of the firm.	Water For Injection manufactured by the same firm of the quoted product.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Valid Certificate of Analysis of the Type / class of glass material used for the vials of the quoted item/s, as issued by the manufacturer of this glass material, coupled with Invoice/proof of purchase: 1. For USP Type 1 glass 4 marks will be awarded. 2. For USP Type 2 Glass 2 marks will be awarded. 3. For product where USP Type 3 glass is used will not be acceptable and will stand disqualified.  Documents duly attested by senior executive of the firm)	Stability studies of quoted item/s not older than 3 years (duly attested by the Q/C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks		
	Ref. No. of item	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	3	3	5	3	3	3	4	4		5	5	5	5	2	3	4	4	5	38	70

Evaluation Criteria for Import of Dry Powdered Injectables 2021-23																											
Name of Firm																											
S. No.	Product General Information				Principal's and Importer's Evaluation Parameters								Suppliers Technical Score	Technical Evaluation Matrix												Product Evaluated Score	Total Technical Score
					Principal Manufacturer Evaluation				Importer's Evaluation					Product Evaluation Parameters													
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25		
					Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 18001/45001 certification	Valid ISO 14001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 14001 certification.	Valid ISO 9001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  Provide Online verification link of ISO 9001 certification	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies/ regulatory body in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificate for equipment in the factory (duly attested by the senior executive of the firm).	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s).  The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.	Adherence to Good Storage Practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC experts at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC experts at the time of inspection).	Valid Proof of export by the Principal Manufacturer to US/Europe/SRA countries/, not older than one year (certificate duly attested by senior executive of the firm).  03 marks for export to SRA Countries. Export Certificate/ COPP/ COMP issued by the relevant regulatory body of the countries mentioned above.	Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	API's source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada, or by regulatory authority of SRAs countries coupled with Form 7 (Batch Certification under Rule 14 D (1)) for awarding marks of API source.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Water For Injection manufactured by the same firm of the quoted product	Valid certificate of the availability of the quoted item in the US market.	Valid WHO prequalification or valid product registration/valid free sale certificate issued by regulatory body of any SRA country(s).	Valid Certificate of Analysis of the Type / class of glass material used for the vials of the quoted item/s, as issued by the manufacturer of this glass material, coupled with Invoice/proof of purchase: 1. For USP Type 1 glass 4 marks will be awarded. 2. For USP Type 2 Glass 2 marks will be awarded. 3. For product where USP Type 3 glass is used will not be acceptable and will stand disqualified.  Documents duly attested by senior executive of the firm)	Stability studies of quoted item/s not older than 3 years (duly attested by the Q/C incharge of the firm).	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health.  Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks					
	Ref. No. of item	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	3	3	5	6	5	6	32	3	5	5	5	2	2	3	4	4	5	38	70		

Evaluation Criteria for Manufacturers of Medical Devices and Sutures for Government MCC 2021-23																								
Name of the firm																								
S.No	Product General Information				Factory Technical Evaluation Parameter								Technical Evaluation Matrix										Product Evaluated Score	Total Technical Score
					Documents Based Factory Score				Evaluation Visit Score				Factory Evaluated Score	Product technical Evaluation Parameters										
	1	2	3	4	5	6	7	8	9	10	11	12		13	14	15	16	17	18	19	20	21		
					Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) <b>Provide verification link for provided for ISO 14001 certification.</b>	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) <b>Provide verification link for provided for ISO 9001 certification.</b>	Valid ISO 13485 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) <b>Provide verification link for provided for ISO 13485 certification.</b>	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Raw material storage (as evaluated at the time of inspection by the MCC expert/s). Non adherence to cGMP shall lead to disqualification of the firm.	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s, shall lead to Disqualification of the firm.	Functional HVAC (as evaluated by the MCC expert/s at the time of inspection). Non-availability or Non functionality of the HVAC system shall lead to Disqualification of the relevant section/firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection. Non-availability of the qualified & relevant HR shall lead to Disqualification of the relevant section/firm).		Current export certificate of the quoted item from DRAP not older than one year (certificate duly attested by senior executive of the firm).  (COPP/COMP or Export NOC issued by DRAP shall be considered) Export to non SRA countries will be awarded (1 mark per country maximum upto 5 marks). Export to any SRA country shall be awarded 5marks.	Goods Declaration certificate of imported raw material of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Raw material Source accredited by WHO, FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by other Regulatory body of SRA countries (Relevant documents duly attested by senior executive of the firm) Coupled with valid proof of purchase of raw Material by the principal manufacturer. In case of API, valid cGMP certificate from DRAP shall be required.	Certificate of Analysis of raw material from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Valid ISO 10993 certificate issued by authorized body of the country of the origin duly accredited with international accreditation forum (IAF) or from IAF accredited body of any SRA country (certificate duly attested by the senior executive of the firm)	Samples evaluation by DTL (Failure to comply with relevant standards shall lead to Disqualification of the quoted products)	Physical examination of the quoted item/s by the MCC expert/s. Rejection of the quoted item/s by the MCC expert/s shall lead to disqualification of the said item/s.				
	Ref. No. of item in MCC Formulary	Generic Name of Item	Trade Name	Size & Gauge of Medical Device	2	3	3	4	2	2	4	5	25	5	5	5	5	3	10	12	45	70		

Evaluation Criteria for Importers of Medical Devices, Government MCC 2021-23																						
Name of the firm																						
S. No.	Product General Parameter				Principal's and Importer's Evaluation Parameters							Technical Evaluation Matrix										
					Principal Manufacturer Evaluation			Importer's Evaluation				Suppliers Technical Score	Product Technical Evaluation									Product Evaluated Score
	1	2	3	4	5	6	7	8	9	10	11		12	13	14	15	16	17	18	19	20	
					Valid ISO 14001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). <b>Provide verification link for ISO 14001 certification.</b>	Valid ISO 9001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). <b>Provide verification link for ISO 9001 certification.</b>	Valid ISO 13485 certificate of the facility where the quoted product is manufactured, issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF) for the country of origin (duly attested by senior executive of the firm). <b>Provide verification link for ISO 13485 certification.</b>	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/regulatory bodies in the case of SRA countries (duly attested by senior executive of the firm)	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s)  <b>The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.</b>	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection).		Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 13, duly attested by the senior executive of the firm.	Raw material Source accredited by WHO, FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by other Regulatory body of SRA countries (Relevant documents duly attested by senior executive of the importer) Coupled with valid proof of purchase of raw Material by the principal manufacturer.	CE/IS (Japanese Free Sale certificate)/US FDA(510 K) certification of the quoted products, 2 marks for each certification, up to a maximum of 06 marks (copies of relevant certificates duly attested by the senior executive of the firm)	Valid ISO 10993 certificate issued by authorized body of the country of the origin duly accredited with international accreditation forum (IAF) or from IAF accredited body of any SRA country (certificate duly attested by the senior executive of the firm)	Samples evaluation by DTL (Failure to comply with relevant standards shall lead to Disqualification of the quoted products)	Physical examination of the quoted item/s by the MCC expert/s. Rejection of the quoted item/s by the MCC expert/s shall lead to disqualification of the said item/s.			
	Ref. No. of item in MCC Formulary	Generic Name of Item	Trade Name	Size, Gauge, etc. of Device	2	3	3	4	4	4	4	24	5	5	5	6	3	10	12	46	70	



Evaluation Criteria for Importers of Cardiac Stents, Government MCC 2021-23																							
	Name of the firm				Technical Evaluation Matrix																		
	Product General Parameters				Principal's & Importer's Evaluation Parameters													Suppliers Technical Score	Product Technical Parameters			Product Evaluated Score	Total Technical Score
					Principal's Evaluation						Importer's Evaluation												
					5	6	7	8	9	10	11	12	13	14	15	16	17						
					Valid cGMP /Quality Control /Quality Assurance Certificate (attested from the embassy of the country of origin in Pakistan or Pakistani embassy in the country of origin). Non provision of the embassy attested certificate will lead to disqualification of firm (duly attested by senior executive of the firm).	Valid certification of US Food and Drug Administration (US FDA) of quoted item/s & Valid permission for sale/import of the quoted item/s in the US market (duly attested by senior executive of the firm). Non Provision of any of these certificates shall lead to disqualification of the quoted item/s.	Valid certificate of accreditation of quoted item/s from European Community (CE) (duly attested by senior executive of the firm).	Valid JIS certification of quoted item/s from Japanese Ministry of Health, Labour & Welfare (MHLW) (duly attested by senior executive of the firm).	Valid ISO 14001 certificate of the facility where the quoted products are manufactured, issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). Provide Online verification link of ISO 14001 certification	Valid ISO 9001 certificate of the facility where the quoted products are manufactured, issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). Provide Online verification link of ISO 19001 certification	Valid ISO 13485 certificate of the facility where the quoted products are manufactured, issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). Provide Online verification link of ISO 13485 certification	Valid accreditation of manufacturing unit or its relevant section/s by International Body (Certificate from US-FDA, WHO and/or other accrediting body /Regulatory bodies from SRA countries duly attested by senior executive of the firm)	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s)  The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.	Adherence to Good storage practices (GSP) for finished goods storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection).		Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 17, duly attested by the senior executive of the firm.	Physical examination of the quoted item/s by the MCC expert/s. Rejection of the quoted item/s by the MCC expert/s shall lead to disqualification of the said item/s.				
	Ref. No. of item in MCC Formulary	Generic Name of Item	Trade Name	Size, Gauge, etc. of Device	5	5	4	4	2	2	3	5	5	5	5	45	5	5	15	25	70		

Evaluation Criteria for Manufacturer of Biological Drugs/Medicines, Government MCC 2021-23																												
Name of Firm:					Technical Evaluation Matrix										Factory Evaluated Score	Product Evaluation Parameters							Product Evaluated Score	Total Technical Score				
					Factory Technical Evaluation Parameters																							
					Documents Based Factory Score					Evaluation visit Score																		
					5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23					
1					2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25
	Ref. No. of Item In	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	2	5	4	4	4	2	5	30	5	5	5	5	5	4	3	3	5			40	70	

Evaluation Criteria for Importers of Biological Medicines/Drugs, Government MCC 2021-23																								
Name																								
				Principal's and Importer's Evaluation Parameters								Suppliers Technical Score	Technical Evaluation Matrix										Product Evaluated Score	Total Technical Score
				Principal Manufacturer Evaluation				Importer's Evaluation					Product Technical Evaluation											
				5	6	7	8	9	10	11	12		13	14	15	16	17	18	19	20	21	22		
				Valid ISO 18001/45001 certificate of the facility where the quoted product is manufactured, issued by authorized bodies/ies of the country of origin duly accredited by International Accreditation Forum (IAF), (duly attested by senior executive of the firm)  (Provide Online verification link of ISO 18001/45001 certification)	Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by authorized bodies/ies of the country of origin duly accredited by International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  (Provide Online verification link of ISO 14001 certification)	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by authorized body of the country of origin duly accredited by International Accreditation Forum (IAF), (duly attested by senior executive of the firm).  (Provide Online verification link of ISO 9001 certification)	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation bodies/ies Regulatory bodies/ies in the case of SRA countries (duly attested by senior executive of the firm)	Valid calibration certificate for equipment in the factory (duly attested by the senior executive of the firm)	Availability of minimum 20% inventory of the total import of the quoted items during last one year (certificate to the effect duly signed by the senior executive of the firm & evaluated by the MCC expert/s). Non availability of the 20 % stock at the time of inspection shall lead to disqualification of the quoted item/s).  The total import detail of the firm during the last one year shall be attested from the Central or concerned Local DRAP office.	Adherence to Good Storage Practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection).		Valid Proof of export by the Principal Manufacturer to US/Europe/SRA countries/ies, not older than one year (certificate duly attested by senior executive of the firm).  03 marks for export to SRA Country/ies.  Export Certificate/ COPP /COMP issued by the relevant regulatory body of the countries mentioned above.	Goods Declaration certificate of imported finished quoted items from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted items, not older than 01 Year on the cutoff date for submission of bids.	API/s source accredited by WHO, US-FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by regulatory authority of SRAs countries coupled with Form 7 (Batch Certification under Rule 14-D (1)) for awarding marks of API source.	Certificate of Analysis of finished quoted items from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Valid Certificate of Analysis of the Type / class of material used for the immediate container of the quoted item/s, as issued by the manufacturer of this material coupled with Invoice/proof of purchase: For award of marks, the certificate of analysis must clearly mention: 1. Pharma Grade Aluminium Foil, PVC, Capsule Shells, Plastic, HDPE or any other material used for the immediate container. 2. Type of Glass material for Liquid ampoules must be USP class 1. 3. Type of Glass material for Oral Syrups/ Suspensions must be USP Type 3 or better. 4. For Dry Powder biologicals USP Type 2	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Stability studies of quoted items not older than 3 years (duly attested by the Q.C incharge of the firm).	Studies on efficacy of products / Biosimilarity Studies on Asian population, as published in PMDC & or HEC recognised journals	Availability of quoted item/s in Pakistani market as per recent most data of IMS/IQVIA Health. Less than 2 % market share = 0 mark 2- 5 % market share = 3 marks More than 5 % market share = 5 marks			
Ref. No. of item in	Generic Name of Item	Dosage Form with Strength	Trade Name	2	2	2	5	5	5	5	4	30	3	5	5	5	4	5	3	5	5	40	70	

Name of the firm					Evaluation Criteria for Manufacturers of Non Drugs Items for Govt MCC 2021-23																									
S.No	Product General Information				Factory Technical Evaluation Parameter												Technical Evaluation Matrix												Product Evaluated Score	Total Technical Score
					Documents Based Factory Score				Evaluation Visit Score								Factory Evaluated Score	Product technical Evaluation Parameters												
					5	6	7	8	9	10	11	12	13	14	15	16		17	18	19	20	21	22							
					Valid ISO 14001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) Provide verification link for ISO 14001 certification.	Valid ISO 9001 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) Provide verification link for ISO 9001 certification.	Valid ISO 13485 certificate of the facility where the quoted product is manufactured, issued by PNAC accredited body (duly attested by senior executive of the firm) Provide verification link for ISO 13485 certification.	Valid accreditation of manufacturing unit or its relevant section/s by the US FDA or WHO or official accreditation bodies regulatory bodies/ies in the case of SRA countries (duly attested by senior executive of the firm)	Raw material storage (as evaluated at the time of inspection by the MCC expert/s). Non adherence to cGMP shall lead to disqualification of the firm.	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s, shall lead to Disqualification of the firm.	Functional HVAC system shall lead to Disqualification of the relevant section/firm.	Adequate availability of qualified & relevant Human Resource (Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection. Non availability of the qualified & relevant HR shall lead to Disqualification of the relevant section/firm).	Current export certificate of the quoted item from DRAP not older than one year (certificate duly attested by senior executive of the firm).  (COPP/COMP or Export NOC issued by DRAP shall be considered) Export to non SRA countries will be awarded (1 mark per country maximum upto 5 marks). Export to any SRA country shall be awarded 5marks.	Goods Declaration certificate of imported raw material of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Declaration of the quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Raw material Source accredited by WHO, FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by other Regulatory body of SRA countries (Relevant documents duly attested by senior executive of the importer) Coupled with valid proof of purchase of raw Material by the principal manufacturer. In case of Pakistani source of API, valid cGMP certificate from DRAP shall be required.	Certificate of Analysis of raw material from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 15, duly attested by the senior executive of the firm.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	Valid ISO 10993 certificate issued by authorized body of the country of the origin duly accredited with international accreditation forum (IAF) or from IAF accredited body of any SRA country (certificate duly attested by the senior executive of the firm)	Physical examination of the quoted item/s by the MCC expert/s. Rejection of the quoted item/s by the MCC expert/s shall lead to disqualification of the said item/s.										
Ref. No. of item in MCC Formulary	Generic Name of Item	Trade Name	Size, Gauge, etc. of Device	2	2	4	5	5	5	5	5	33	5	5	5	5	2	3	12	37	70									

Evaluation Criteria for Importers of Non-Drug Items (NDIs) Government MCC 2021-23																							
S. No.					Technical Evaluation Matrix																	Product Evaluated Score	Total Technical Score
					Principal's and Importer's Evaluation Parameters							Suppliers Technical Score	Product Technical Evaluation										
					Principal Manufacturer Evaluation				Importer's Evaluation														
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21			
				Valid ISO 14001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). Provide verification link for ISO 14001 certification.	Valid ISO 9001 certificate of the facility where the quoted product is manufactured issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF), (duly attested by senior executive of the firm). Provide verification link for ISO 9001 certification.	Valid ISO 13485 certificate of the facility where the quoted product is manufactured, issued by authorized body of the country of origin duly accredited with International Accreditation Forum (IAF) for the country of origin (duly attested by senior executive of the firm). Provide verification link for ISO 13485 certification.	Valid accreditation of manufacturing unit or its relevant section/s by the US-FDA or WHO or official accreditation body/ies/regulatory body/ies in the case of SRA countries (duly attested by senior executive of the firm)	Availability of minimum 20% inventory of the total import of the quoted item/s during last one year of certificate to the effect (certificate signed by the senior executive of the firm & evaluated by the MCC expert/s at the time of inspection shall lead to disqualification of the quoted item/s).  The total import detail of the firm during the last one year shall be attested by the central or concerned Local DRAP office.	Adherence to Good storage practices (GSP) for finished good storage of the quoted item/s. Non adherence to GSP, as evaluated by the MCC expert/s at the time of inspection shall lead to Disqualification of the firm.	Adequate availability of qualified & relevant Human Resource Certified by the senior executive of the firm & evaluated by MCC expert/s at the time of inspection).		Goods Declaration certificate of imported finished quoted item/s from Pakistan Customs, coupled with valid airway bill or Bill of Lading for the quoted item/s, not older than 01 Year on the cutoff date for submission of bids.	Raw material Source accredited by WHO, FDA, EMA, MHRA, TGA, PMDA, Swiss Medic or Health Canada or by other Regulatory body of SRA countries (Relevant documents duly attested by senior executive of the importer) Coupled with valid proof of purchase of raw Material by the principal manufacturer.	Certificate of Analysis of finished quoted item/s from the Principal Manufacturer as mentioned in the goods declaration (GD) provided in column 13, duly attested by the senior executive of the firm.	Valid WHO prequalification or valid product registration / valid free sale certificate issued by regulatory body of any SRA country(s).	CE/JIS (Japanese Free Sale Certificate) / US FDA 510-K) certification of quoted item/s, 01 mark for each of the listed certification, up to a maximum of total 03 marks (copies of relevant certificates duly attested by the senior executive of the firm).	Valid ISO 10993 certificate issued by authorized body of the country of the origin duly accredited with international accreditation forum (IAF) or from IAF SRA country (certificate duly attested by the senior executive of the firm)	Physical examination of the quoted item/s by the MCC expert/s. Rejection of the quoted item/s by the MCC expert/s shall lead to disqualification of the said item/s.					
	Ref. No. of item in MCC Formulary	Generic Name of Item	Trade Name	Size, Gauge, etc. of Device	2	2	4	5	5	6	6	30	5	5	5	3	3	3	16	40	70		

## Section VI. Sample Forms

### **MANDATORY STANDARD FORMS (1 to 5)**

**BID FORM 1: BID COVER SHEET**

**BID FORM 2: LETTER OF INTENTION**

**BID FORM 3: AFFIDAVIT**

**BID FORM 4: PRICE SCHEDULE FORMAT FOR FINANCIAL BID**

(To be submitted in separate sealed envelope)

**BID FORM 5: INTEGRITY PACT**

**CONTRACT AGREEMENT** (for information only, shall be signed by the successful bidders only)

# **Bid Form-1**

## **BID COVER SHEET**

### **Mandatory General Information of Applicant Firm**

**NOTE: Complete filling of this form along with the provision of all requisite information is mandatory. Missing or not providing any of the requisite information may lead to disqualification of the bidder/s from the bidding competition without any correspondence. Any appeal from bidder/s, for whatsoever reasons, shall not be entertained in such a case.**

S.No.	Name of the Bidding Firm:	
<b>1.</b>	Please indicate whether the firm is: <ul style="list-style-type: none"> <li>i. Manufacturer, or</li> <li>ii. Importer, or</li> <li>iii. Both; Manufacturer as well as Importer</li> </ul> For various MCC formulary items offered for this bidding competition.	
<b>2.</b>	Please indicate out of the following category/ies, under which the Firm is applying for bidding: <ul style="list-style-type: none"> <li>i. General medicines</li> <li>ii. I/V Fluids</li> <li>iii. Biological drugs</li> <li>iv. Medical devices including Surgical Disposables, Cotton &amp; related goods, gauze, adhesive tapes, bandages, etc., but excluding cardiac stents</li> <li>v. Cardiac Stents</li> <li>vi. Non drug items (NDIs).</li> </ul>	
<b>3.</b>	Please provide names, attested copies of CNICs, two recent attested photographs, valid street addresses in Pakistan, all working landline, mobile phone numbers and valid email address of the following: <ul style="list-style-type: none"> <li>i. Owner/Proprietor of the Firm; and</li> <li>ii. Managing Director / CEO of the Firm; and</li> <li>iii. Focal person shall be an employee of the firm/bidder officially authorized for day to day official correspondence/communication if required with the procuring agency along with valid mobile number.</li> </ul> 2. Please provide clear, legible and visible attested photocopies of all the valid requisite items mentioned items)	
<b>4.</b>	Please provide the following valid information regarding applicant Firm: <ul style="list-style-type: none"> <li>i. Complete street address of the:             <ul style="list-style-type: none"> <li>a. Head Office</li> <li>b. Main warehouse; and</li> </ul> </li> <li>ii. Valid &amp; working official Landline Phone and Fax Numbers; and</li> <li>iii. Valid Mobile phone number/s of the Focal Person registered which should be registered his/her CNIC No. and name; and</li> <li>iv. Valid and functional Email address; and</li> <li>v. Official Website address/es.</li> </ul>	

5.	<p><b>i.</b> Please provide, in original, the bids security instrument amounting to Rupees Eight Hundred Thousands only (Rs.800,000/-) in the shape of Call Deposit Receipt (CDR) in the name of the Director General Health Services, Khyber Pakhtunkhwa, along with the Financial Proposal in the sealed envelope, from a scheduled Bank of Pakistan. Ordinary crossed or open Cheques shall not be acceptable as Bids security.</p> <p><b>ii.</b> Note: Please also provide an attested photocopy of the same bids security document in the sealed envelope of technical Proposal.</p>
6.	<p>Please provide attested copies of the following Tax related valid documents:</p> <p><b>iii.</b> National Tax Number (NTN) of the Firm for Income Tax, and</p> <p><b>iv.</b> Last year Income Tax Return of the Firm; and</p> <p><b>v.</b> Sale Tax Registration Certificate of the Firm; and</p> <p><b>vi.</b> Certificate of Professional Tax of the Firm.</p>
7.	<p>In case of being a Manufacturer, the Firm should provide attested copies of the following documents also:</p> <p><b>i.</b> Valid Drugs Manufacturing License issued by the Drugs Regulatory Authority of Pakistan (DRAP); and</p> <p><b>ii.</b> Valid Product Registration Certificate issued by the DRAP for the item/s quoted by the Firm for this bidding competition</p> <p><b>iii.</b> Valid cGMP certificate issued by DRAP</p> <p><b>iv.</b> Valid Price List of the quoted item/s</p> <p><b>v.</b> Dissolution Profile for each quoted drug / medicine item belonging to the category of oral dosage form.</p>
8.	<p>In case of being Importers, the Firm should provide attested copies of the following documents also:</p> <p><b>i.</b> Valid Drugs Sales License for the importer; and</p> <p><b>ii.</b> Valid Product Registration Certificate issued by the DRAP for the imported item/s quoted by the Firm for this bidding competition; and</p> <p><b>iii.</b> Valid Agency Agreement with the Foreign Principal Manufacturer entity/ies; and</p> <p><b>iv.</b> Valid cGMP Certificate/ Quality Control /Quality Assurance Certificate / Certificate of Pharmaceutical Products (COPP)/Certificate of Medicinal Products (COMP) of the Principal Manufacturer from an IAF accredited body or any accredited / authorized / regulatory body in the country of origin for the quoted item/s as issued for the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm; (In case of Non-applicability of the above mentioned certificates for Examination Gloves (Non Sterile) and Adhesive Tapes (Non Sterile) only, provision of EC-Declaration of conformity from the principal manufacturer (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s is mandatory) and</p> <p><b>v.</b> Valid Free Sale Certificate for the quoted item/s as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s). Non provision of this document shall lead to disqualification of the firm; and</p> <p><b>vi.</b> Valid Price List of the quoted items.</p> <p><b>vii.</b> Establishment of Medical Device License issued by DRAP for the item/s quoted by the firm for bidding competition.</p> <p><b>viii.</b> For cardiac stents, provision of the following documents is mandatory apart from those mentioned in clause a &amp; b above:</p> <p><b>i.</b> Valid US-FDA certificate of the quoted item/s; and</p> <p><b>ii.</b> Valid permission of sale or import of quoted item/s for sale in the US open market.</p> <p>Note: Valid cGMP, COPP and COMP certificate/s, and Valid Free Sale Certificate/s for the quoted item/s, as issued by relevant authority of the country of origin of the quoted imported good/s (duly attested from the Embassy / High Commission / Consulate (as the case may be) of the country of origin in Pakistan or Pakistani Embassy / High Commission / Consulate (as the case may be) in the country of origin of the quoted good/s), as elaborated in the relevant section of these SBDs, shall be presented in original by the bidder to the inspection team of MCC expert/s at the time of inspection . Failure to comply with this instructions shall lead to disqualification of the firm for the quoted item/s. Photocopy or scanned copy of the same shall not be considered in lieu of the original.</p>

9.	<p>The bidding Firm shall also provide an Affidavit on Judicial Stamp Paper of the value of at least Rs. 100/- (Rs. One Hundred Only) for the following undertaking:</p> <ul style="list-style-type: none"> <li><b>i.</b> I / We have carefully read the whole set of Standard Bidding Documents for this bidding competition and that I / We have fully understood and agree to all the provisions (including, but not limited to, those provided under ITB 11.5, 16.1 and 29.1 of the Bid Data Sheet), terms and conditions, evaluation criteria, mechanism of evaluation &amp; selection of items for which the Firm has applied for competition; and</li> <li><b>ii.</b> I / We fully understand and agree that the bidding competition for which I / We have applied to enter in, shall be based on merit based scoring system for the evaluation of technical bids which has inverse relationship with</li> </ul>
	<p>the rates quoted by the bidders in their financial bids submitted; and that in this situation, the lowest financial bid/s may or may not win the bidding competition; and</p> <ul style="list-style-type: none"> <li><b>iii.</b> I / We guarantee that the quoted drug / medicine items are, and shall be, freely available in the market of Pakistan; and particularly in the market of Khyber Pakhtunkhwa Province; and</li> <li><b>iv.</b> I / We shall provide to the inspection team/s of expert/s authorized for the purpose by the Directorate General Health Services Khyber Pakhtunkhwa; an uninterrupted and free access to all relevant documents, sections of the manufacturing facilities / unit, storage and warehousing facilities as well as any other area relevant, as deemed appropriate by the above mentioned team for their purpose of visit/s.</li> <li><b>v.</b> In case any documents submitted in relation to this bidding competition or any undertaking given by the Firm, if found incorrect or false or misleading or diverting the decision making for the competition, shall be liable to be proceeded for blacklisting for any business with / by the Government of Khyber Pakhtunkhwa, Health Department, confiscation of bids security and / or any other lawful action as deemed appropriate by the Government of Khyber Pakhtunkhwa, including that to be taken in concert with the DRAP or any other body / entity of the Federal Government; and</li> <li><b>vi.</b> I / We have fully understood that the medical devices and items in the categories of cotton, bandages, adhesive tapes, etc. including other non-drug items shall be evaluated / examined by expert/s nominated by the Technical Evaluation Committee / Selection &amp; Rate Contracting Committee of the Government MCC of the Health Department, Khyber Pakhtunkhwa at its sole discretion; and that the Firm shall fully agree and abide by the decision / opinion, whatsoever, of the said expert/s regarding the selection, or otherwise, of the quoted item/s for purchase / rate contracting.</li> <li><b>vii.</b> I / We also undertake that submission of any false/bogus/fake/forged/ fabricated/tampered document shall lead to disqualification of our firm from this bidding competition as well as to other lawful action/s to be taken by the concerned authorities.</li> <li><b>viii.</b> I / We have fully understood that no such documents shall be entertained by the Procuring Agency, which is issued after due date of Bid opening.</li> </ul>
10.	<p>I certify and affirm that I have attached /provided all the requisite mandatory documents / information including Bids Security with this Bid and that I fully understand that any document if not provided / missing shall result in the disqualification and declaring my bid as ineligible and thus non-responsive.</p> <p>Signatures: _____</p> <p>Name: _____</p> <p>CNIC No. _____</p> <p>Designation: _____</p> <p>Address: _____</p>

## **Bid Form-2**

### **Letter of Intention**

*Bid Ref No.*

*Date of the Opening of Bids*

*Name of the Contract :{ Add name, e.g, Supply of Dugs and Medicines, etc.}*

*To: [Name and address of Procuring Agency]*

Dear Sir/Madam

Having examined the bidding documents, including Addenda Nos. *[insert numbers& Date of individual Addendum]*, the receipt of which is hereby acknowledged, we, the undersigned, offer to supply and deliver the Goods under the above-named Contract in full conformity with the said bidding documents and at the rates/unit prices described in the financial bid are not more than the trade price of quoted item/s in the market.

We undertake, if our bid is accepted, to deliver the Goods in accordance with terms and condition of contract agreement.

We agree to abide by this bid, for the Bid Validity Period specified in the Bid Data Sheet and it shall remain binding upon us and may be accepted by you at any time before the expiration of that period.

Until the formal final Contract is prepared and executed between us, this bid, together with your written acceptance of the bid and your notification of award, shall constitute a binding Contract between us.

We understand that you are not bound to accept the lowest or any bid you may receive.

We undertake that, in competing for (and, if the award is made to us, in executing) the above contract, we will strictly observe the laws against fraud and corruption in force in Pakistan.

Dated this *[insert: number]* day of *[insert: month]*, *[insert: year]*.

Signed:

In the capacity of *[insert: title or position]*  
Duly authorized to sign this bid for and on behalf of *[insert: name of Bidder]*

## **Bid Form-3**

### **AFFIDAVIT** *(on Judicial Stamp Paper)*

I/We, the undersigned [**Name of the Supplier**] hereby solemnly declare and undertake that:

- 1) I / We, the undersigned, have read the contents of the Bidding Document and have fully understood it.
- 2) The Bid being submitted by the undersigned complies with the requirements enunciated in the bidding documents.
- 3) The Goods that I / We, the undersigned, propose to supply under this contract are eligible goods within the meaning of this SBD.
- 4) The undersigned are also eligible Bidders within the meaning of the Standard Bidding Documents.
- 5) The undersigned are solvent and competent to undertake the subject contract under the Laws of Pakistan.
- 6) The undersigned have not paid nor have agreed to pay, any Commissions or Gratuities to any official or agent related to this bid or award or contract.
- 7) The undersigned are not blacklisted or facing debarment from any Government, or its organization or project.
- 8) That undersigned has not employed any child labor in the organization/unit.
- 9) We understand that the Procuring Agency or any of its committees are not bound to accept the lowest or any other bid they may receive.

I / We affirm that the contents of this affidavit are correct to the best of my/our knowledge and belief.

Signatures with stamp

Name: \_\_\_\_\_

Designation: \_\_\_\_\_

CNIC No. \_\_\_\_\_

For Messrs. [**Name of Supplier**]

## **Bid Form-4**

**Note:** *This form is to be submitted in a separate sealed envelope to be kept within the main sealed envelope of the bid.*

### **Price Schedule format for Financial Bid of Government MCC for the year 2021-23**

- 1. In case of Drugs/Medicines,** the unit price of each item shall be quoted and submitted in the following format:

S.No.	Serial No. of quoted Drug / Medicine in the MCC Formulary 2021-23	Generic Name with Strength and Dosage Form of quoted Drug / Medicine	Trade/Brand Name of quoted Drug / Medicine	Maximum Retail Price (MRP) of the quoted items	Trade Price of quoted Drug / Medicine (Unit price)	Rate Offered per unit in Pak. Rupees (Rs) for quoted Drugs / Medicines.
1						

**Note:** Quoted price of the items shall be rounded up to two decimal points. For Example, Rs. 16.34/.

- 2. In case of Surgical Disposables, Medical Devices (Type 1 and 2) (NDIs),** the unit price of each item shall be quoted and submitted in the following format:

S.No.	Serial No. of quoted item in the MCC Formulary 2021-23	Generic Name with sizes/measurements of quoted item	Trade / Brand Name of quoted item	Maximum Retail Price (MRP) of the quoted item	Trade Price of quoted item (Unit price)	Rate Offered per unit in Pak. Rupees (Rs) for the quoted item
1						

**Note:** Quoted price of the items shall be rounded up to two decimal points. For Example, Rs. 16.34/-

## **Bid Form-5**

**INTEGRITY PACT** (on Judicial Stamp Paper)

**Declaration of Fees, Commission and Brokerage Etc. Payable by Suppliers of Drugs/Medicines,  
Surgical Disposables, Medical Devices & Non Drugs Items for Govt: MCC 2021-23**

In response to advertisement related to the bidding process / competition regarding purchase and supply of drugs, non-drugs and surgical disposable items for 2021-23 for the health facilities / institutions through directorate general Health Services, Khyber Pakhtunkhwa, Peshawar, I, Mr. / Ms. \_\_\_\_\_ s/o, d/o \_\_\_\_\_ bearing CNIC No. \_\_\_\_\_, and having the Designation of \_\_\_\_\_ in Messrs.

(M/S) [***Name of Supplier***] do hereby solemnly affirm, declare and certify on behalf of M/S [***Name of Supplier***] that:

1. [***Name of Supplier***] has not obtained or induced the procurement of any contract, right, interest, privilege or other obligation or benefit from Government of Khyber Pakhtunkhwa (GoKP) or any administrative subdivision or agency thereof or any other entity owned or controlled by GoKP through any corrupt business practice; and
2. That without limiting the generality of the foregoing, [***Name of Supplier***] represents and warrants that it has fully declared the brokerage, commission, fees etc. paid or payable to anyone and not given or agreed to give and shall not give or agree to give to anyone within or outside Pakistan either directly or indirectly through any natural or juridical person, including its affiliate, agent, associate, broker, consultant, director, promoter, shareholder, sponsor or subsidiary, any commission, gratification, bribe, finder's fee or kickback, whether described as consultation fee or otherwise, with the object of obtaining or inducing the procurement of a contract, right, interest, privilege or other obligation or benefit in whatsoever form from GoKP, except that which has been expressly declared pursuant hereto; and
3. That [***Name of Supplier***] has made and will make full disclosure of all agreements and arrangements with all persons in respect of or related to the transaction with GoKP and has not taken any action or will not take any action to circumvent the above declaration, representation or warranty; and
4. That [***Name of Supplier***] accepts full responsibility and strict liability for making any false declaration, not making full disclosure, misrepresenting facts or taking any action likely to defeat the purpose of this declaration, representation and warranty. It agrees that any contract, right, interest, privilege or other obligation or benefit obtained or procured as aforesaid shall, without prejudice to any other rights and remedies available to GoKP under any law, contract or other instrument, be voidable at the option of GoKP; and
5. That notwithstanding any rights and remedies exercised by GoKP in this regard, [***Name of Supplier***] agrees to indemnify GoKP for any loss or damage incurred by it on account of its corrupt business practices and further pay compensation to GoKP in an amount equivalent to ten times the sum of any commission, gratification, bribe, finder's fee or kickback given by [name of Supplier] as aforesaid for the purpose of obtaining or inducing the procurement of any contract, right, interest, privilege or other obligation or benefit in whatsoever form from GoKP.

**Signatures with stamp**

**Name:** \_\_\_\_\_

**Designation:** \_\_\_\_\_

**CNIC No.** \_\_\_\_\_

**For Messrs. [***Name of Supplier***]**

**Witness No. 1**

**Witness No. 2**

(Signatures, name, father's name, CNIC & address of each Witness)

## **Bid Form-6**

### **GOVERNMENT MCC RATE CONTRACT AGREEMENT (for successful bidders)**

**THIS RATE CONTRACT AGREEMENT** is made and agreed today on the \_\_\_\_ day of [Month], 2021 between the Director General Health Services, Health Department, Government of Khyber Pakhtunkhwa (*hereinafter referred to as the Procuring Agency or first party, which expression shall, where the context admits, be deemed to include the successors and / or assignee/s of the Provincial Government of Khyber Pakhtunkhwa*); and Messrs. [Name of Supplier] through Mr.

\_\_\_\_\_  
Designation

\_\_\_\_\_  
CNIC No. \_\_\_\_\_, (*hereinafter referred to as the Supplier or second party or he or his or him, which expression, unless repugnant to the context, means and includes their legal heir/s, successors-in-interest, assignee/s and legal representative/s*) that:

**WHEREAS** the Procuring Agency has made a bidding competition under the approved Standard Bidding Documents for the year 2021-23 (*hereinafter referred to as the SBDs*) approved for the selection and rate contracting of drugs/medicine, medical devices, surgical disposables and other non-drug items (*hereinafter referred to as goods*) for actual purchases of the selected and rate contracted goods to be made by the offices / officers of the Health Department, Government of Khyber Pakhtunkhwa (*hereinafter called the Purchasing Agency or Purchasing Agencies or Purchasing Agency/ies, where the context so admits*); and

**WHEREAS** the Supplier has won the bidding competition for selected goods, as listed in the Schedule-1 of this contract agreement; and

**WHEREAS** the Supplier declares that he is not a broker, middle-man, distributor or authorized dealer of, or acting on behalf of any entity or person, but himself a genuine Manufacturer and / or direct Importer of the goods for which he has won the bidding competition for supply of the same to the Purchasing Agency/ies, as defined in the SBDs, throughout the province of Khyber Pakhtunkhwa (*hereinafter referred to as the Province*); and

**WHEREAS** both the parties have agreed that the Purchasing Agencies in the Province shall purchase all, or some, or none of the goods, as of details given in the Schedule-1 of this Contract Agreement, from the Supplier at the sole discretion of the individual Purchasing Agency/ies in subordination to laws and matters ancillary to the terms and conditions of the SBDs; and

**WHEREAS** the Supplier shall supply all the goods ordered by the Purchasing Agency/ies to the latter in the quantity as mentioned in the supply order to be issued by the Purchasing Agency within the timeframe as mentioned in clause-22 of this contract agreement;

**Now, therefore,** both the parties hereby mutually agreed to enter into this contract agreement as under:

1. The Supplier agrees to take full responsibility of the validity and implications, that may arise in future, of declaration as submitted by him through an affidavit on judicial stamp paper along with the Bid Form-1 of the SBDs along with his bid; and also that in case of any kind of breach of the said declaration, the Supplier shall be liable to be proceeded against by the Procuring Agency and / or Purchasing Agency concerned, as the case may be, in accordance with the clauses of this rate contract agreement as well as relevant laws, rules and regulations of the Government of Khyber Pakhtunkhwa, as amended from time to time, to govern the situation/s.
2. The Supplier shall supply the ordered goods to the concerned Purchasing Agency exactly at the address of the official premises situated within the district of the official jurisdiction of the latter as provided in the supply order issued to the former.
3. The Supplier shall be solely responsible for the safe and appropriate method and mode of transportation, loading, unloading and staking of the supplied items till, and at the time of delivery to the destination address indicated by the Purchasing Agency in the district of its jurisdiction.

4. The Supplier shall be solely responsible for any damage, untoward incidence, maintenance of required temperature and protection from light and other environmental conditions as well as other hazards that may possibly or potentially affect the safety, quality and efficacy of the supplied goods till the time of delivery and the consequences arising therefrom, if any.
5. The Supplier shall not claim or charge any transportation, loading / unloading, labour or any other charges, whatsoever, related to or in the name of logistics, accidents, insurance, freight, toll tax, etc.
6. The Supplier shall supply all the goods in full conformity to the specifications as laid down in the SBDs.
7. The Purchasing Agency shall arrange to obtain randomized sample/s for each formulary item of the supplied goods, as in the SBDs and belonging to the categories of drug/medicine, medical devices and surgical disposables through the notified Drug Inspector/s concerned for sending the same to the concerned Drug Testing Laboratory for Test / Analysis as provided in the Drugs Act 1976, DRAP Act 2012 and rules frame thereunder as well as provisions of the SBDs, further subject to the following condition/s:
  - a. The supplied goods declared in contravention to any provision of the Drugs Act 1976, DRAP Act 2012 and rules made thereunder, shall be replaced by the Supplier at his sole risk and cost and at no cost to the Purchasing Agency, within 07 days from the date of intimation to the Supplier or his focal person, as nominated by the Supplier in the Bid Form-1 of his bid submitted under the SBDs, at such place as the Purchasing Agency may direct in accordance with clause-2 of this contract agreement.
  - b. The Purchasing Agency shall arrange to obtain sample/s of the replaced goods as in clause-7 (a) above, for the purpose of Test / Analysis as provided in the Drugs Act 1976, DRAP Act 2012 and rules made thereunder.
  - c. In case of non-supply or delayed supply or partial supply of replacement items, as in clause-7 (a) above, the Supplier shall be liable for imposition of one or more penalties as provided in clause-22 of this contract agreement.
  - d. All the contravened stock of goods, as in clause-7(a) above, if seized by the authorities shall be the case property under the provisions contained in the Drugs Act, 1976 and the rules made thereunder.
  - e. The supplier shall be responsible to make arrangements for appropriate storage and the matters ancillary to the safe custody of the seized case property as in clause-7(d) above at his sole risk, cost and responsibility with no claim, whatsoever, from the concerned Purchasing Agency, and / or the Drug Inspector, and / or Procuring Agency. The firm will also produce batch wise cold chain data from the source of origin & thermoLog data from factory to ware house for temperature sensitive drugs.
  - f. In case the destruction of the seized stock, as in clause-7(d),(e) above, is required to be undertaken under the applicable laws and rules, all the costs involved in the execution of the decision and destruction, whatsoever, shall be solely borne by the supplier without any claim of any nature, whatsoever, from the concerned Purchasing Agency or Drug Inspector or Procuring Agency.
  - g. Any of the item, as per clause-7 above, if initially declared to be in contravention with any provision of Drugs Act 1976, but later on declared as of standard quality by the concerned Appellate Drugs Testing Laboratory, shall be returned to the supplier by the concerned Drug Inspector in a lawful manner.
8. Supplier shall supply to the Purchasing Agency the freshly manufactured goods having maximum possible long expiry dates with the minimum remaining shelf life of at least 65% in case of imported goods and at least 85% in case of locally manufactured goods within Pakistan.

9. The Supplier shall hoist the list of supplied goods on his official website, while indicating name of items, name of manufacturer / importer, Invoice No., warranty & date, Registration No., Batch No., quantity, unit price and expiry date of the supplied goods along with the name of the Purchasing Agency.
10. In case of taking any action in contravention to any provision of the applicable law and rules, the Supplier shall render himself liable to such lawful action/s as deemed appropriate and taken against him under any or all the applicable law/s, rule/s of the Government of Khyber Pakhtunkhwa, terms and conditions of the SBDs and the clauses of this contract agreement.
11. The Purchasing Agency shall recommend to the Procuring Agency for taking legal / lawful action against the Supplier regarding non-supply, short supply, substituted supply, delayed supply or any other unlawful action / shortcoming, on the part of Supplier, pertaining to the Drugs Act 1976 or in the execution of this contract agreement.
12. The Procuring Agency shall take lawful / legal action against the Supplier in accordance with the clauses of this contract agreement as well as relevant and applicable laws, rules and regulations of the Government of Khyber Pakhtunkhwa, as amended from time to time, to govern suchlike situation/s, which may, inter alia, include but not limited to blacklisting, forfeiture of earnest money and performance guarantee, if any.
13. The Supplier agrees to the following conditions related to packing, packaging and labelling of the goods to be supplied to Purchasing Agencies under this contract agreement:
  - a. Each item shall be supplied to Purchasing Agency in the packing and packaging unit as approved and registered by the DRAP. The supplier shall supply all the unit items bearing the words **"GOVERNMENT OF KHYBER PAKHTUNKHWA MCC SUPPLY"** and **"NOT FOR SALE"** in block letters and clearly visible manner with indelible ink, **along with the name of the Purchasing Agency concerned** on the label, outer packing of each individual unit item as well as on its outer carton/s.
  - b. The labels shall comply with all the requirements as laid down under the Drugs Labelling and Packing Rules 1986. The strip / blister shall clearly indicate expiry date of the same medicine in a clear and legible manner.
  - c. The goods shall be packed and transported to the Purchasing Agency in accordance with the provisions contained in the Standard Bidding Documents.
  - d. The items related to the category of Absorbent Cotton / Surgical Gauze / Cotton Bandages / Crepe bandage, etc. shall be supplied in strict compliance with the instructions contained in Notification No. F.6-6/2005-Reg-II (south) dated 13/9/2006 of the then Federal Ministry of Health, Pakistan.
14. The Procuring Agency or its representative shall have the right to inspect the manufacturing facilities, premises, warehouses, godowns, laboratories etc. at any time during the financial year 2021-23 /or till the execution of supply orders given under this contract agreement by the Purchasing Agency of the Province. If anything found in contravention of cGMP, clauses of Drugs Act 1976 or of this Contract Agreement the Procuring Agency shall have the sole right and authority to take any lawful action as deemed appropriate, against the Supplier which may include, but not limited to cancellation of supply order/ orders given to the Supplier by the Purchasing Agency as well as imposition of penalties, forfeiture of supplied stock, forfeiture of performance guarantee or earnest money as the case may be, stoppage or recovery of payment made to the supplier as well as taking any other lawful action.
15. The Supplier agrees that the approved price of all individual items in Schedule-1 of this contract agreement, as quoted by him in the financial bid, shall remain valid till and up to 30<sup>th</sup> June 2023.

16. As mentioned in Special Conditions of Contract, the bid security of Rs. 800,000/- from the Supplier as already received by the Procuring Agency at the time of bids submission under GCC Clause 15, shall be retained by the Procuring Agency as Performance Security till the end of contract period and will be released back to supplier in response to applying for the same by him to the Procuring Agency after successful completion of all the contractual obligations of this contract agreement and the SBDs.
17. The Supplier shall provide legal and valid warranty to the Purchasing Agency for all the goods supplied under this contract agreement, which fall under the provisions of Drugs Act 1976, DRAP Act 2012 and the rules framed thereunder, on prescribed Form-2A in accordance with the mechanism prescribed for the purpose.
18. For Non-Drug Items, the Supplier shall provide appropriate warranty to the Purchasing Agency in accordance with Special Conditions of Contract of the SBDs for this bidding competition, for each item supplied in response to supply orders.
19. In case the Supplier had been awarded marks during the technical evaluation for API source accreditation for Drugs / Medicines, and for medical grade material certification for medical devices & Non-Drug Items, and for Pharmaceutical grade certification for immediate containers of Drugs/medicines shall warranty the supply of all such goods with the same certified quality, material and specification to the Purchasing Agency throughout the validity period of this contract agreement.
20. Bill for payment in triplicate along with all other relevant and required documents shall be submitted by the Supplier to the Purchasing Agency immediately after completion of supply of ordered stock. The Supplier shall be bound to pay all sorts of government taxes, duties and stamp duties, imposed earlier or during the financial year by the Government of Pakistan or by the Provincial Government of Khyber Pakhtunkhwa on any supplied / purchased item.
21. In case of situation related to Force Majeure, the Supplier may immediately without delay inform the Procuring Agency as well as the Purchasing Agency in writing about the situation along with solid proof of the situation through the fastest, lawful and available means of communication, but not through the electronic mail, and request the Procuring Agency for the grant of extension in the supply period.
  - a. The Procuring Agency, in case of being fully satisfied with the genuineness of situation arising from the claimed Force Majeure by the Supplier, may extend the period of supply of goods up to a maximum of not more than thirty days.
  - b. The Procuring Agency and / or Purchasing Agency shall, in no case, be responsible or held responsible for any complications in making payments to Supplier by the Purchasing Agency that may arise from the closure of financial year, and / or lapse, and / or surrender of public funds, vis-à-vis, the standard and normal public sector financial management laws, rules, regulations, procedures and practices governing the Procuring Agency, and / or Purchasing agency/ies.
  - c. After the expiry of extended period as in clause-21(a) above, the supply order shall stand cancelled to the extent of non-supplied goods and the performance security in the form of retained bids security, as in clause-16 of this contract agreement shall be forfeited in favour of the Procuring Agency.
22. The Supplier agrees that the supply of the ordered goods under this agreement shall be completed by the Supplier i.e. Local Manufacturer within thirty (30) days and Importer Supplier within sixty (60) days after the receipt of supply order/s from the Purchasing Agency/ies, except in situation/s covered under clause-21 above regarding Force Majeure. In case of delay in supplies reaching to the Purchasing Agency, the following penalties shall be imposed by the Purchasing Agency upon the Supplier:

- a. Upon delay in supply beyond 30 and 60 days for local manufacturer supplier and for importer supplier respectively a lump sum penalty of 1% per week shall be deducted upto a maximum of 7% penalty for 7 weeks, of the total quoted price of such goods, whose supply was delayed out of the same supply order as issued to the supplier, shall be levied through deducting the total amount of penalty from the total pre-tax payable billed amount by the Purchasing Agency.
- b. In case of delay in supply beyond 7 weeks after the cutoff days, as mentioned in clause-22 above, the supply order issued by the Purchasing Agency shall stand cancelled to the extent of non-supplied items and in such a case, the Procuring Agency shall have the right, duty and authority to impose any or all of the below mentioned penalties; that is
  - i. Forfeiting the bids security and / or performance guarantee of the Supplier as related to this contract agreement; and / or
  - ii. Immediately debarring the selected item/s and/or Supplier/firm from future participation and business not less than one year and up to next three (03) calendar years with the Government of Khyber Pakhtunkhwa through MCC or any other health institution, project and / or Program directly or indirectly run or implemented by or through the provincial Health Department or Purchasing Agencies in the Province, as defined in the SBDs, and District Governments in the Province; and / or
  - iii. Initiating the process for and recommending for permanent blacklisting of the Supplier with the Purchasing Agencies under Debarment/Blacklisting Guidelines Notified vide Letter No. 2440-2500/Proc. Cell, Dated: 30-08-2018.
  - iv. The applicant bidder shall be debarring from the process of contract frame work agreement 2021-23 either for its quoted item/s and/or firm from the bidding competition at any stage where he has been declared defaulter firm/non supplied firm in the contract period of 2021-23 (**30<sup>th</sup> June 2023**) reported by purchasing agencies as a non-supplier firm and proceeded by procuring entity as per Debarment/Blacklisting Guidelines of Health Department.
23. Notwithstanding any rights, duties and / or remedial measures and / or managerial actions taken and / or to be taken and / or any powers exercised and / or to be exercised by the Procuring Agency and / or Purchasing Agency and / or Purchasing Officer/s with regard to the execution of this contract agreement, the Supplier agrees to indemnify all of them for any loss or damage incurred or inflicted upon by them in individual or official capacity upon the Supplier whether through any of their actions and / or practices and / or otherwise.
24. The Supplier further agrees to pay compensation to the Government of Khyber Pakhtunkhwa of an amount equivalent to ten times the sum of any commission, gratification, bribe or kickback and / or finder's fee given by the Supplier for the purpose of obtaining and / or inducing the procurement of any contract, right, interest, privilege or other obligation/s or benefit/s in whatsoever form, from the Procuring Agency or any of the Purchasing Agencies.
25. The supplier further agrees that all the data related to supplies throughout the financial year shall be provided to the procuring entity by the end of financial year. The CDR of the supplier shall not be released till the provision of the said data.
26. The Procuring Agency and / or Purchasing Agency, as the case may be, and the Supplier shall make every effort to resolve amicably by direct negotiation any disagreement or dispute arising between them under or in connection with the contract / supplies. However, despite such negotiation if the Purchasing Agency & Supplier have been unable to resolve amicably a contract dispute, either party may refer the case to Secretary to Government of Khyber Pakhtunkhwa, Health Department, Peshawar for decision through a Dispute Resolution Committee under the chairmanship of Special Secretary Health with Additional Secretary

Health (Development) or Additional Secretary Health (Establishment) and Deputy Secretary Drugs as members.

27. Both the parties agree that the Procuring Agency in the capacity of being the overall head of the Government Medicine Coordination Cell, or otherwise, has the authority to regulate, if deemed appropriate, under the provisions in the SBDs, through imposing restrictions and / or classifying and / or grouping any selected quoted item/s for stopping, increasing or decreasing the purchase of such item/s by the Purchasing Agency/ies to rationalize and / or control the use and / or misuse of such item/s.

---

**Director General Health Services  
Khyber Pakhtunkhwa  
For and on behalf of Government of  
Khyber Pakhtunkhwa,  
Health Department, Peshawar**

---

Signature:  
Name:  
Designation  
CNIC No.  
Stamp:

**For and on behalf of Manufacturers /  
Importer**

**WITNESS NO. 1  
Chief Pharmacist  
Government MCC, DGHS  
Health Department, Khyber  
Pakhtunkhwa, Peshawar**

**WITNESS NO. 2  
Signature:  
Name:  
Father's Name:  
Address:  
CNIC No.**

### **Schedule -1**

**Directorate General Health Services, Khyber Pakhtunkhwa**

**Government MCC 2021-23**

1. **Name and Address of Supplier:**
2. **List of Selected Item/s from the Supplier along with quoted unit price/s:**