TENDER REFERENCE NO.: DRG/92/2026/PHARM(TC)

MINISTRY OF HEALTH
NEGARA BRUNEI DARUSSALAM

SUPPLY AND DELIVERY OF MEDICINES FOR THE DEPARTMENT OF PHARMACEUTICAL SERVICES FOR A PERIOD OF THREE (3) YEARS

TENDER FEES : $100.00

RECEIPT NO. :

CLOSING DATE : 21st April 2026

TIME : 2.00 PM

FOA :

THE CHAIRMAN
MINI TENDER BOARD, TENDER BOX
GROUND FLOOR, MINISTRY OF HEALTH
COMMONWEALTH DRIVE
BANDAR SERI BEGAWAN BB3910
NEGARA BRUNEI DARUSSALAM

(NON-CLUSTERING)

SECTION 2

SPECIFICATIONS

TENDER REFERENCE NO.: DRG/92/2026/PHARM(TC)

INVITATION TO TENDER
SUPPLY AND DELIVERY OF MEDICINES FOR THE DEPARTMENT OF PHARMACEUTICAL SERVICES MINISTRY OF HEALTH FOR A PERIOD OF THREE (3) YEARS

NO.	ITEM DESCRIPTION	ESTIMATED ANNUAL REQUIREMENTS	BUFFER STOCK (UNITS)	PACKING / PRESENTATION	PACK SIZE	DELIVERY PERIOD
1	Imatinib (as mesilate) 400mg oral solid preparation preferably tablet	130 x 30's	20% of Estimated Annual Usage	Blister pack with 10's per strip preferred. Preference will be given to suppliers quoting item where the name and strength of the product appears over each blister pocket or be oriented centrally	-	First order within 2 months upon receipt of purchase order, subsequent order 1 month

Please note that only medicinal products registered with BDMCA will be considered. In cases where the active ingredient is not registered with BDMCA for certain reasons, products registered in other countries, such as those listed as BDMCA-benchmark countries, may then be considered, subject to case-to-case evaluation.

The following documents and/or information are required with each offer. Failure to comply with the requirement may cause unnecessary delay in processing for approval from the relevant authority

NO.	REQUIREMENTS	ENTER RESPONSE HERE
1	Validity of offer price shall be at least 12 months from the closing date of submission of quote.	
2	**Sample** Vendor is required to submit sample in untampered original pack including package insert and Summary of Product Characteristics (in English) For Controlled drugs and Psychotropics: In the event that sample cannot be submitted, photos and/or artwork are required. See 'Presentation'.	
3	**Presentation** Vendor is to submit: i. Clear colour-printed photo of the product offered with company's official stamp. Photo must show label details of the primary and secondary packaging from all sides/angles including name / brand of item, strength and form / preparation. ii. High resolution photo of the following ▪ For tablets / capsules: ✓ Appearance of individual tablets / capsules; ✓ If the item is in strip pack, the back and front of the strip ▪ For Injections: ✓ Appearance of individual vial / ampoule / syringe	
4	**Registration with Brunei Darussalam Medicines Control Authority (BDMCA)** A copy of any of the following: ▪ Product Licence Certificate ▪ Letter of authorization from product licence holder, if applicable Priority will be given to medicinal products already registered with the BDMCA.	
5	**Manufacturer details** Please provide manufacturer details with supporting documents. If manufacturer details are not available, please provide the following: i. Hardcopy of declaration letter from the principal to inform any issues related to product safety, quality and efficacy as a result of a recall by the manufacturer, wholesaler or Department of Health ii. A copy of the principal's wholesaler license.	
6	**Shelf life** Please indicate the product shelf-life. Priority will be given to product with a minimum of 24 months.	
7	**Storage Condition** The storage labelling should be in accordance with the latest guideline on registration of medicinal products in Brunei Darussalam. Priority is given to products with specified storage conditions. Terms such as "ambient conditions", "room	

NO.	REQUIREMENTS	ENTER RESPONSE HERE
	temperature" or "does not require any special storage condition" should be avoided unless stability studies are provided.	
8	**Alcohol and animal content** Declaration of source of animal origin and/or alcohol content (if any) is to be provided.	
9	**Certificate of Analysis** A copy of the product's Certificate of Analysis (CoA) is to be submitted.	
10	**New Product** Where the product offered has never been supplied to the Ministry of Health, Brunei, detailed information on the product is to be submitted. The information required include, but not limited to, the following: i. Bioequivalence studies (Generic products) and / or Clinical studies ii. Stability studies iii. Certificate of free sales iv. Certificate of Pharmaceutical Product (CPP) v. A copy of the Summary of Product Characteristics vi. Good Manufacturing Practice (GMP) certificate vii. Batch release certificate or certificate of origin (for blood products)	
11	**Price Justification** Vendor is to submit letter of justification on price increase if the same product has been previous supplied to Ministry of Health from the same supplier / distributor.	
12	**Local content & Tax Compliance Certificate** Vendor is to provide a copy of the latest content of the company as well as the updated tax compliance certificate, if applicable	
13	**Product Registration Number in any of the *benchmark/reference countries** If not registered with BDMCA, priority will be given to products which are registered in any of the benchmark/reference countries. *****The benchmark/reference countries are Australia, Canada, Malaysia, Singapore, United Kingdom, the United States of America, France, Sweden, Japan, Switzerland, Republic of Korea & European Union.	
14	**Patent Declaration** i. Vendors quoting generic products must provide an official declaration or confirmation from the Brunei's Intellectual Property Office verifying whether the innovator product is off-patent ii. Vendors quoting innovator products must provide information on the patent status and expiry in Brunei.	

Preference will be given to medicinal products already:

- **Registered with the BDMCA.**

SECTION 3

FORM TO BE USED

CONTENTS

SCHEDULE 1

TENDER FORM

To:

TENDER REFERENCE NO.: DRG/92/2026/PHARM(TC)

INVITATION TO TENDER
SUPPLY AND DELIVERY OF MEDICINES FOR THE DEPARTMENT OF PHARMACEUTICAL SERVICES MINISTRY OF HEALTH FOR A PERIOD OF THREE (3) YEARS

TENDER OF *(name of tenderer)* _____

Company/Business Registration No _____

Tender Closing Date: _____

NO.	ITEM	BRAND NAME	MANUFACTURER	PACKING/ PRESENTATION	PACK SIZE	UNIT PRICE (B$)	TOTAL PRICE (B$)
1	Imatinib (as mesilate) 400mg oral solid preparation preferably tablet						
DELIVERY PERIOD: First order 2 months upon receipt of purchase order, subsequent order 1 month							
DELAY AND LIQUIDATED DAMAGES: If the Supplier fails or is unable to deliver the Goods or any parts thereof on the Delivery Date within the time specified, the Government shall be entitled, without prejudice to claim from the Supplier by way of liquidated damages for each day of such delay, a sum of equal to one percent (1%) of the price of the Goods as stated in the relevant Purchase Order in respect of the delayed delivery, provided that the total liquidated damages shall not exceed ten percent (10%) the Purchase Order Price.							

1. We offer and undertake on your acceptance of our Tender to supply and deliver the above-mentioned goods in accordance with your Invitation To Tender.

2. Our Tender is fully consistent with and does not contradict or derogate from anything in your Invitation To Tender. We have not qualified or changed any of the provisions of your Invitation To Tender.

3. OUR OFFER IS VALID FOR **TWELVE (12)** CALENDER MONTHS FROM THE TENDER CLOSING DATE. Where the price validity period differs from that required by the Government i.e. 12 months, the **LONGER VALIDITY PERIOD** will be taken as the final validity period.

4. When requested by you, we shall extend the validity of this offer.

5. We further undertake to give you any further information which you may require.

Dated this_____day of_____

…… Tenderer's official stamp:
Signature of authorised officer of Tenderer
Name:
Designation:

SCHEDULE 2 - INFORMATION SUMMARY

2.1 Tenderers shall provide in this Schedule the following information:

a. Management summary

b. Company profile (including Contractor and sub-contractor(s), if any)

c. Years of experience (as of the Tender Closing Date) of the Contractor and sub-contractor(s) in the:
   - *Supply and Delivery of Medicines.*

d. Other information which is considered relevant

SCHEDULE 3 – SUB-CONTRACTS

3.1 Tenderers shall complete Table 3.1 with information about all the companies involved in the provision of the services and items specified in this tender. This shall include details about the Contractor and each sub-contractor involved, as well as their respective responsibilities.

3.2 Tenderers shall also indicate in Table 3.1 any alliance relationship established with each sub-contractor. An alliance is defined as a formal and binding business relationship between the allied parties.

Table 3.1 - Responsibility Table

Company Name	Responsibility Description	Alliance Relationship between Contractor and Sub-contractor(s)		
		Alliance Exists? (Y/N)	Date Established	Alliance Description
Contractor				
		Not Applicable	Not Applicable	Not Applicable
Sub-contractor(s)				

SCHEDULE 4 – COMPANY'S BACKGROUND

4.1 Each of the companies involved in this tender, including Contractor and sub-contractor(s) (if any), shall provide information on the company's background, scope of operations, financial standing and certified copy of its Certificate of Incorporation or Certificate of Registration (as the case may be).

SCHEDULE 5 – REFERENCES

5.1 Tenderers shall submit a list of customers in Table 5.1 to whom the Contractor has provided similar services and items as specified in this tender in the recent 5 years as of the Tender Closing Date.

Table 5.1 - References of previous customers

Customer Name and Address	Customer Type (Govt or Quasi Govt) *	Contact Person	Title	Contact Number, Fax Number and E-mail Address

***Note: Tenderers shall indicate whether the customer is a Government or Quasi Government organisation. A Quasi Government is defined as an organisation which (1) is managed and controlled by the Government; or (2) has at least 50% shares being held by the Government. Please leave the column blank if the customer is neither a Government or Quasi Government organisation.**

5.2 The Ministry of Health shall treat all the information submitted under this schedule in strict confidence.

5.3 The Ministry of Health reserves the right to contact the references for tender assessment purposes.

SCHEDULE 6 - SUBMISSION OF SAMPLE

6.1 Tenderers shall submit the Submission of Sample form below in respect of the items specified in this tender.

6.2 Samples of the items to be submitted shall be:

a) identical in packing and manufacture to the items to be offered by the Tenderer; and

b) marked with the corresponding item number of the tender.

SUBMISSION OF SAMPLE FORM

To:

TENDER REFERENCE NO: DRG/92/2026/PHARM(TC)

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SUBMISSION OF SAMPLE FORM OF (NAME OF TENDERER)

NO.	ITEM	SAMPLE SUBMITTED (indicate ✓)	SAMPLE NOT SUBMITTED (indicate x with a reason)
1	Imatinib (as mesilate) 400mg oral solid preparation preferably tablet		

We understand as stated in the Instructions To Tenderers that Tenders without samples shall not be considered unless they meet the criteria for sample exemptions.

…….
(*Signature of authorized officer of Tenderer*)
Name:
Designation:
Date:

Tenderer's official stamp:

FOR OFFICE USE

Date of receipt: _____

Receiving Officer: _____

SCHEDULE 7

LETTER OF DECLARATION

SCHEDULE 8

PRICE JUSTIFICATION FORM

TENDER REFERENCE NO.: DRG/92/2026/PHARM(TC)

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SUBMISSION OF PRICE JUSTIFICATION LETTER OF (NAME OF TENDERER)

Please attach price justification letter if brand offered in this tender had been supplied previously to the Ministry of Health and has increased in price.

NO.	ITEM	Price justification letter submitted (Indicate ✓ or X or Not offered)	Letter reference (if applicable)
1	Imatinib (as mesilate) 400mg oral solid preparation preferably tablet		

Dated this_____day of _____

…… Tenderer's official stamp:
Signature of authorised officer of Tenderer
Name:
Designation: