KENYA MEDICAL SUPPLIES AUTHORITY

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KENYA MEDICAL SUPPLIES AUTHORITY — YOUR PARTNER IN HEALTHCARE

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All Correspondence should be addressed to Chief Executive Officer

When replying please quote our ref:

To: All Prospective bidders, **Date: 24th March, 2023**

CLARIFICATION 1

Dear Sir/Madam,

The following questions were raised by interested bidders and responses provided.

Question 1

We are requesting change of Pack size for Item no: PM10MPL004-Misoprostol 200 Mcg Tablets (sublingual) Pack of 28s to Pack of 30s;

i. For fair competition since only one supplier has the Pack of 28s as compared to Pack of 30s.

ii. MSK's packaging is packs of 20's. Please advise.

Response

Note change of pack size to:

Misoprostol 200 mcg Tablets (sublingual) Pack of 30s

Question 2

The Mifepristone +Misoprostol should be a pack of 30's. However, MSK's packaging is for 50's. Please advise.

Response

The required product and pack size is:

Mifepristone 200mg+Misoprostol 200 mcg- pack of 30's

Question 3

Is it compulsory to have PPB approved manufacturing facility and PPB Market Authorization for the product to participate in the tender?

Response

The question touches on facility and product and therefore here are the requirements:

Current and valid Good Manufacturing Practice (GMP) certificate issued by an independent body or Market Authorization issued by the Pharmacy and Poisons Board of Kenya. *(This is specific to a manufacturing plant/site/facility)*

For products registered within the year, provide Product Registration certificate issued by the Pharmacy and Poisons Board of Kenya. For products registered in prior years, provide Product Registration certificate and or Retention Certificate with QR codes issued by the Pharmacy and Poisons Board of Kenya. *(This is specific to a product).*

Question 4

We have a plant that has PPB-GMP but the products are yet not registered at PPB. Can we participate in the below Tenders?

KEMSA/OIT07-2022-2024

KEMSA-OIT08-2022-2024

KEMSA-OIT09-2022-2024

KEMSA-OIT10-2022-2024

Response

With regards to product registration and retention, the criteria below shall apply.

For products registered within the year, provide Product Registration certificate issued by the Pharmacy and Poisons Board of Kenya. For products registered in prior years, provide Product Registration certificate and or Retention Certificate with QR codes issued by the Pharmacy and Poisons Board of Kenya.

Question 5

Kindly let us know If the forms mentioned below need to be included in the document: -FORM ELI 1.1, FORM FIN 3.1, FORM FIN 3.2, FORM CON 1 , FORM EXP 1 & FORM PER 1. If yes, kindly provide the template.

Response

Bidders should fill in the templates that are mandatory under specific evaluation criteria.

Question 6

Product PM10ETP005 ERYTHROPOETIN 2000IU/0.5ML (VIAL) is what is needed in the specification. However, if ours is 2000IU/0.5ml (Pre filled Syringe) can we still quote for it?

Response

Yes, we shall require erythropoietin 2000iu/0.5mL pre filled syringe and **NOT** a vial.

Question 7

My Hydralazine strength is 20mg/ml ampoule in a box pack of 5. Please clarify if the strength is ok?

Response

The required strength is 20mg/mL.

Question 8

Please let us know if the below forms need to be provided as they have not been included in the tender document. If yes, kindly provide the templates.

Form ELI-1.1- Tenderer information form

Form FIN-3.1- Financial situation & performance

Form FIN 3.2- Average annual turnover

Form CON-1- Current contract commitments/ contracts in progress form

Form EXP-1- Experience

Form PER-1- Historical contract non-performance, and pending litigation and litigation history

Response

Bidders should fill in the templates that are mandatory under specific evaluation criteria.

Yours faithfully

Dr. Justus Kinoti(Ph.D)
Ag. DIRECTOR,PROCUREMENT
FOR:CHIEF EXECUTIVE OFFICER