

2024/07/16

تاريخ إنشاء العطاء:

P37-2024-H

اسم العطاء :

الصلاحية:

& Reagents  
Consumables For  
Hepatitis Tests And  
Blood Unit Sreening

وصف العطاء:

الوثيقة المالية:

طريقة الشراء:

Item No	Stock No	Stock Name	Quantity	Unit
1	09_6_00_008	Hepatitis C Virus Antibody/Automated approved methods	123600	Test
2	09_6_00_009	Human Immune Deficiency Virus 1& 2 Antibody/Automated approved methods	115200	Test
3	09_6_00_010	Hepatitis B Surface Antigen /Automated approved methods	123600	Test
4	09_6_00_011	Hepatitis B Surface Antibody/Automated approved methods	9000	Test
5	09_6_00_012	Hepatitis B Envelope Antigen/ Automated approved methods	1200	Test
6	09_6_00_013	Hepatitis B Envelope Antibody/ Automated approved methods	1200	Test
7	09_6_00_014	Hepatitis B Core Total Antibody/ Automated approved methods	98400	Test
8	09_6_00_015	Hepatitis B Core Antibody IgM/ Automated approved methods	900	Test
9	09_6_00_016	Hepatitis A Virus Antibody IgM/ Automated approved methods	7200	Test
10	09_6_00_017	Syphilis ( Treponema Antibody Detection Kit)/ Automated approved methods	97200	Test
11	09_6_03_022	HAV-IgG Antibodies By ELISA: Detection of total antibodies against HAV in human serum of plasma.	8160	Test

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Item No	Stock No	Stock Name	Quantity	Unit
12	09_6_03_023	Hepatitis C Virus (HCV) By ELISA in human serum or plasma	28800	Test
13	09_6_03_028	HIV-Ab 1+2 by ELISA,	28800	Test
14	09_6_03_034	Hepatitis A-IgM-Ab kit,by ELISA	960	Test
15	09_6_03_038	Hepat B Surface Ag Kit, by ELISA	28800	Test
16	09_6_03_045	Hepatitis Delta Ab by Elisa	96	Test
17	09_6_03_0451	Hepatitis Delta Ag by Elisa	96	Test
18	09_6_03_046	Hepatitis Kit, Hepatitis B Core Ab Total IgG by Elisa	28800	Test
19	09_6_03_047	Hepatitis B Surface Ab Kit,ELISA HBs-Ab	576	Test
20	09_6_03_049	HCV Recombinant Immunoblot Assay.	72	Test
21	09_6_06_0622	HEV IgM antibody for Hepatitis E by Elisa	96	Test

شروط خاصة

- 1-Number of shipments will be decided by purchasing committee according to the respective minimum guaranteed shelf life upon delivery/dispatch to be provided in the original offers to establish delivery schedule accordingly .The DRMS with this respect reserves the right to reject any item not in compliance with this term and impose the correspondent fines.
- 2-Dispatched/Delivered good must show Description of goods , expiry date must be clearly indicated on all package and relative documents ,Batch No., Storage conditions, stamped "Sold to RMS" ,and any other necessary information .
- 3- goods must be dispatched/delivered under the same storage conditions that comply with their nature.
- 4-offers to include all kit size configurations from Manufacturer and number of tests per kit/pack to be clearly mentioned where applicable.
- 5-All packages to be stamped or labelled with the following "Sold to RMS" Tender P37-2024-H".
- 6-Any Products derived or containing materials of human origin should be supplied with a certificate and/or official document from the manufacturer stating that the products are devoid of blood borne infectious agents :HbsAg, HIV, HCV, ect. Items belonging to this category to be specified in the original offer.
- 7- Goods should be previously evaluated and approved in DRMS or purchased by "central procurement branch " or by " directorate of defence procurement ".
- 8-The DRMS reserves the right to purchase goods in respect to their kit size (tests/kit) which will correlate with technical needs of the specific department.

9-Offers must include a full warranty including spare parts and labour for a period of minimum of 24 months from the date of first shipment for all the existing machine.

10-Any additional requirements for test performance should be supplied as a free of charge according to the number of tests (such as Lamps, reaction cells sets, sample probe substrate, calibrator, control, sample cups, sample cuvettes, cuvettes probe wash, cleaning solutions, wash solutions, multi diluents, universal diluents, disposable pipett tips, IgG absorbent) must be mentioned clearly, and the company must provide us with prices list of these free material.

11-A copy of (CE or FDA certificated) is to be submitted and should be certificated by JFDA (Jordan food & drug administration).

12- Custom clearance of goods shall be the responsibility of the Jordanian Armed Forces (JAF), however, suppliers shall bear all costs incurred by handling charges and any demurrage charges or extra expenses incurred by the port's corporation or QAIA (including expenses caused by delay in presenting the necessary shipment documents for either clearing or transporting the goods to the required location mentioned in the final order, delivery note issuing charges, unloading charges, local shipping charges, etc. ). The supplier is also responsible for providing of all relevant shipping documents, together with the delivery order(s).

13- Pricing must include services of sale, shipment, transportation, delivery from port to site or Main Medical Stores.

14- For offers submitted in Jordanian dinars payment will be either by wire transfer or by cheque to be paid after receiving DOCUMENTS FROM MAIN MEDICAL STORES, any other way of payment will not be accepted and the offers will be rejected by the purchase committee.

15-All items should retain at least the MSL (8) months for (MSL). Any item not complying with this term upon receipt should be accompanied with a confirmation that you accept to replace any remaining unused quantities at your expenses after the expiry date. Said confirmation is subject to the approval of the director of the Royal Medical Services and will incur a fine, which will be decided later by DRMS according to the loss that this discrepancy with terms of the tender has caused.

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16-All necessary reagents for the ELIZA Technique kits (96 well plate , controls , calibrations , standards ,conjugate ,TMB or substrate solution ,diluent ,buffers ,wash solutions ,sulfuric acid or stop solution ) should be packed within the kit (complete kit) as whole unit in the country of origin.

17-Bidders must submit their reservation /queries regarding tender specificareqtions and/or special term within the first half of the tender closing period starting from the tender announcement date . Reservation /queries submitted after the end of this period shall be rejected.

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