الإعلان

الى : السادة

م/مناقصة عامة لشراء المستلزمات الطبية

رقم: SUP 98/2026/19R على الموازنة الجارية

رقم كتاب الدعوة:

يسر وزارة الصحة/ الشركة العامة لتسويق الادوية والمستلزمات الطبية (كيماديا) بدعوة مقدمي العطاءات المؤهلين و ذوي الخبرة لتقديم عطاءاتهم لتجهيز **(مستلزمات اكياس دم / نبيذة اساسي وكما في الجدول المرفق)** مع ملاحظة ما يأتي: .

Sup 98 -2026-19 R - (CIP)

NO.	NATIONAL CODE	ITEM	UOM	الملاحظات	QTY.	الكلفة التخمينية	منشأ الكلفة النخمينية
1	DIS-DE00-035	Single plastic blood bag disp. sterile with CPDA1 anticoagulant solution 450ml capacity	PCS	"-Product name: Single blood bag with sampling arm system - Description: Single bag CPDA1-450ml collection and for storage whole blood (mandatory). The system must include sampling device with vacuum holder and needle protector (for after collection) - Blood bag volume: 450/ ml(mandatory) -Needle and needle cap: Size of needle "16G" with needle cap (mandatory) Once the cap opens, must be tamper-evident that the cap has been opened and not able to lock again Needle system should have indicator for bevel up. (mandatory) - Needle protection: The needle protection is welded(mandatory). This avoids occasional opening and guarantees the product integrity a) Once the blood collection is completed, needle must be smoothly pulled by the operator into the needle protector device (NPD) and lock(mandatory). Must be signaled to personnel by an audible click or tactile indication. b) When the collection needle is locked in the NPD, the entrance surface of NPD must extend at least 8mm beyond the tip of the needle (to be evaluated). The collection needle should not slide out from the NPD c) After use, the NPD should interlock with the vacuum tube holder. (to be evaluated) d) the needle with less pain (to be evaluated) - Clamp: Clamp is used for connecting off blood flow. -Sampling system: The sampling tube and donor tube must have clamps. The sampling system is completely assembled ready to accept vacuum sampler(mandatory). The system is designed in order to avoid interference with the blood flow and to reduce the risk of activation of the coagulation factors. Sampling pouch: Capacity of 30 - 50mL (to be evaluated). Vacuum tube holder: The barrel should be transparent (mandatory), and the barrel must extend at least 18mm beyond the tip of the sampling needle (to be evaluated). The break valve should be placed after the Y connector to the donor tube (and not on the sampling pouch tube) (to be evaluated) to prevents leakage form the donor needle. -Break-off connectors: The break-off cones position is forced to let the blood components easily flow in both directions(mandatory) With audible click. -"Y" connectors: The ((Y)) connectors have a reduce inner volume to avoid turbulence to the blood components flow. - Labels(mandatory) Complying with ISO standard 3826. Batch number and product identification code are reported also and / or in bar code system according to the ISBT standard. All, main labels must be tamper-proof Graphical symbol of the type of blood component Liable for addition stickers -Packing: (mandatory) Protective dual packing (individual and (Plastic or aluminum) Complying with ISO Standard 3826 . - Certifications should be supplied by manufacturer's company FDA or CE MARKED - Plastic film: Medical grad PVC (class VI) complying with ISO specification 3826 (should be supplied by manufacturer's company) - Anticoagulant:(mandatory) CPDA1 63ml. - Validity: (mandatory) 24 month or more - Indications: For collection and storage of whole blood"&	251057	3.23	اوربي

2	DIS-DE00-036	Quadruple plastic Blood bag sterile with CPDA1 solution 450ml capacity	PCS	"- Product name: Quadruple plastic blood bag with CPDA1 solution 450ml - Description: (Name and /or Graphical symbol) Quadruple bag CPDA1 for preparation and storage of products (plasma, red cell and platelet and cryo. Precipitate) with blood sampling arm system adaptor for sampling with vacuum containers and with protector needle system (for needle protection after collection) (mandatory) - Blood Bag volume: Four bags 450 ml with CPDA1 63ml, 400 ml platelet gas permeable and (400ml, 400 ml) plasma bags (mandatory) - Needle: Size of needle "16G" with needle's cap(mandatory). Once the cap opens, must be tamper-evident that the cap has been opened and not able to lock again. Needle system should have indicator for bevel up(mandatory) - Needle protection: The needle protection is welded(mandatory). This avoids occasional opening and guarantees the product integrity a) Once the blood collection is completed, needle must be smoothly pulled by the operator into the needle protector device (NPD) and lock(mandatory). Must be signaled to personnel by an audible click or tactile indication. b) When the collection needle is locked in the NPD, the entrance surface of NPD must extend at least 8mm beyond the tip of the needle (to be evaluated). The collection needle should not slide out from the NPD. c) After use, the NPD should interlock with the vacuum tube holder. (to be evaluated) - Clamp: Clamp is used for connecting off blood flow. The sampling tube and donor tube must have clamps(mandatory). - Sampling system: The sampling system is completely assembled ready to accept vacuum sampler(mandatory). The system is designed in order to avoid interference with the blood flow and to reduce the risk of activation of the coagulation factors. Sampling pouch: Capacity of 30 - 50mL (to be evaluated) Vacuum tube holder: The barrel should be transparent (mandatory), and the barrel must extend at least 18mm beyond the tip of the sampling needle (to be evaluated). The break valve should be placed after the Y connector to the donor tube (and not on the sampling pouch tube) (to be evaluated) to prevents leakage form the donor needle. - Break-off connectors: The break-off cones position is forced to let the blood components easily flow in both directions (mandatory) With audible click. -"Y" connectors (mandatory): The ((Y)) connectors have a reduce inner volume to avoid turbulence to the blood components flow. - Labels:(mandatory) Complying with ISO standard 3826. Batch number and product identification code are reported also and / or in bar code system according to the ISBT standard. All, main labels must be tamper-proof 6 Name and / or Graphical symbol of the type of blood component on each bag Liable for addition stickers - Packing: (mandatory) Protective dual packing (individual and (Plastic or aluminum) Complying with ISO Standard 3826. - Certifications should be supplied by manufacturer's company FDA or CE MARKED -Plastic film : Medical grad PVC (class VI) complying with ISO specification 3826 (should be supplied by manufacturer's company) - Anticoagulant(mandatory). CPDA1 63ml - Validity(mandatory) 24 month or more - Indications:(mandatory) For collection of whole blood, preparation and storage plasma, red blood cell and platelet and cryo.precipitate"&	369312	8.5	اوربي

3	DIS-DE00-037	Quadruple plastic Blood bag sterile with CPD /SAGM solution 450ml capacity (with leukocyte filter)	PCS	"- Product name: Quadruple blood bag with filter for leukodepletion of RBC. - Description:(Name and / or Graphical symbol) Quadruple bag CPD/SAGM in line filter for leukodepletion of RBC, blood sampling arm system adaptor for sampling with vacuum containers and with protector needle system (for needle protection after collection). (mandatory) - Blood Bags volume: Four bags 450 with CPD 63ml, 400 ml for platelets concentrate and gas permeable),(400ml plasma and 400 ml (with 100ml SAGM solution) (mandatory) - Needle: Size of needle "16G" with needle's cap.(mandatory) Once the cap opens, must be tamper-evident that the cap has been opened and not able to lock again. Needle system should have indicator for bevel up(mandatory). - Needle protection: The needle protection is welded(mandatory). This avoids occasional opening and guarantees the product integrity a) Once the blood collection is completed, needle must be smoothly pulled by the operator into the needle protector device (NPD) and lock(mandatory). Must be signaled to personnel by an audible click or tactile indication. b) When the collection needle is locked in the NPD, the entrance surface of NPD must extend at least 8mm beyond the tip of the needle (to be evaluated). The collection needle should not slide out from the NPD. c) After use, the NPD should interlock with the vacuum tube holder. (to be evaluated) - Clamp: Clamp is used for connecting off blood flow. The sampling tube and donor tube must have clamps(mandatory). -Sampling system: The sampling system is completely assembled ready to accept vacuum sample(mandatory). The system is designed in order to avoid interference with the blood flow and to reduce the risk of activation of the coagulation factors. Sampling pouch: Capacity of 30 - 50mL (to be evaluated). Vacuum tube holder: The barrel should be transparent(mandatory), and the barrel must extend at least 18mm beyond the tip of the sampling needle (to be evaluated). The break valve should be placed after the Y connector to the donor tube (and not on the sampling pouch tube) (to be evaluated) (to prevents leakage form the donor needle. - Break-off connectors: The break-off cones position is forced to let the blood components easily flow in both directions. (mandatory) With audible click -"Y" connectors: The ((Y)) connectors have a reduce inner volume to avoid turbulence to the blood components flow. (mandatory) - Labels(mandatory) Complying with ISO standard 3826. Batch number and product identification code are reported also and / or in bar code system according to the ISBT standard. All, main labels must be tamper-proof Name and / or Graphical symbol of the type of blood component on each bag Liable for addition stickers - Packing: (mandatory) Protective dual packing (individual and (Plastic or aluminum) Complying with ISO Standard 3826 - Filter: Filter for leukocyte removal from RBC. - Filtering matter: - Red cell recover > 90% (to be evaluated) -Certifications should be supplied by manufacturer's company FDA or CE MARKED - Plastic film: Medical grad PVC (class VI) complying with ISO specification 3826 (should be supplied by manufacturer's company) - Anticoagulant:-(mandatory) CPD 63ml / SAGM 100ml - Validity(mandatory) 24 month or more. - Indications: Preparation of: Concentrated and filtered red blood cell CPD solution and SAGM solution is used for Storage red blood cell for 42 days and Plasma and platelets concentrate."&	456131	19.98	اوربي

4	DIS-DE00-038	Double plastic Blood bag sterile with CPDA1 solution 450ml capacity	PCS	"- Product name: Double blood bag with sampling arm syste - Description:(Name and / or Graphical symbol) Double bag CPDA1-450ml for preparation and storage of red blood cell and PRP products with blood sampling arm system adaptor for sampling with vacuum containers and with protector needle system (for needle protection after collection) (mandatory) - Blood Bag volume(mandatory) Two bags (450ml and 350 ml) - Needle: Size of needle "16G" with needle's cap.(mandatory) Once the cap opens, must be tamper-evident that the cap has been opened and not able to lock again. (mandatory) Needle system should have indicator for bevel up(mandatory) - Needle protection: The needle protection is welded(mandatory). This avoids occasional opening and guarantees the product integrity a) Once the blood collection is completed, needle must be smoothly pulled by the operator into the needle protector device (NPD) and lock(mandatory). Must be signaled to personnel by an audible click or tactile indication. b) When the collection needle is locked in the NPD, the entrance surface of NPD must extend at least 8mm beyond the tip of the needle. The collection needle should not slide out from the NPD. (to be evaluated) c) After use, the NPD should interlock with the vacuum tube holder. (to be evaluated) - Clamp: Clamp is used for connecting off blood flow. The sampling tube and donor tube must have clamps. (mandatory) - Sampling system: The sampling system is completely assembled ready to accept vacuum sampler. (mandatory) The system is designed in order to avoid interference with the blood flow and to reduce the risk of activation of the coagulation factors. Sampling pouch: Capacity of 30 - 50mL (to be evaluated). . Vacuum tube holder: The barrel should be transparent(mandatory), and the barrel must extend at least 18mm beyond the tip of the sampling needle .(to be evaluated) The break valve should be placed after the Y connector to the donor tube (and not on the sampling pouch tube) (to be evaluated () to prevents leakage form the donor needle. - Break-off connectors: The break-off cones position is forced to let the blood components easily flow in both directions. (mandatory) With audible click -"Y" connectors(mandatory) The ((Y)) connectors have a reduce inner volume to avoid turbulence to the blood components flow. - Labels(mandatory) Complying with ISO standard 3826. Batch number and product identification code are reported also and / or in bar code system according to the ISBT standard. All, main labels must be tamper-proof Name and / or Graphical symbol of the type of blood component on each bag Liable for addition stickers - Packing(mandatory) Protective dual packing (individual and (Plastic or aluminum) Complying with ISO Standard 3826 . -Certifications should be supplied by manufacturer's company FDA or CE MARKED -Plastic film: Medical grad PVC (class VI) complying with ISO specification 3826 (should be supplied by manufacturer's company) - Anticoagulant. (mandatory) CPDA1 63ml. - Validity(mandatory) 24 month or more . - Indications:(mandatory) For collection of whole blood, preparation and storage of red blood cell and PRP products"&	150736	4.85	اوربي

5	DIS-DE00-039	Triple plastic Blood bag sterile with CPDA1 solution 450ml capacity	PCS	"-Product name: Triple blood bag with sampling arm system - Description: (Name and / or Graphical symbol) Triple bag CPDA1-450ml for preparation and storage of products plasma, red cell and platelet platelets concentrate with blood sampling arm system adaptor for sampling with vacuum containers and with protector needle system (for needle protection after) collection(mandatory) - Blood Bag volume: Three bags (450ml, 400 ml platelets concentrate and gas permeable) and 350 ml plasma) (mandatory) - Needle: Size of needle "16G" with needle's cap.(mandatory) Once the cap opens, must be tamper-evident that the cap has been opened and not able to lock again. Needle system should have indicator for bevel up(mandatory) - Needle protection: The needle protection is welded(mandatory). This avoids occasional opening and guarantees the product integrity a) Once the blood collection is completed, needle must be smoothly pulled by the operator into the needle protector device (NPD) and lock(mandatory). Must be signaled to personnel by an audible click or tactile indication. b) When the collection needle is locked in the NPD, the entrance surface of NPD must extend at least 8mm beyond the tip of the needle. The collection needle should not slide out from the NPD(to be evaluated). c) After use, the NPD should interlock with the vacuum tube holder (to be evaluated) -Clamp: Clamp is used for connecting off blood flow. The sampling tube and donor tube must have clamps. (mandatory) - Sampling system: The sampling system is completely assembled ready to accept vacuum sampler. (mandatory) The system is designed in order to avoid interference with the blood flow and to reduce the risk of activation of the coagulation factors. Sampling pouch: Capacity of 30 - 50mL (to be evaluated). Vacuum tube holder: The barrel should be transparent(mandatory), and the barrel must extend at least 18mm beyond the tip of the sampling needle. (to be evaluated) The break valve should be placed after the Y connector to the donor tube (and not on the sampling pouch tube) (to be evaluated) to prevents leakage form the donor needle. - Break-off connectors: The break-off cones position is forced to let the blood components easily flow in both directions(mandatory) With audible click. 14 -"Y" connectors(mandatory): The ((Y)) connectors have a reduce inner volume to avoid turbulence to the blood components flow. - Labels(mandatory) Complying with ISO standard 3826. Batch number and product identification code are reported also and / or in bar code system according to the ISBT standard. All, main labels must be tamper-proof Name and / or Graphical symbol of the type of blood component on each bag Liable for addition stickers - Packing(mandatory) Protective dual packing (individual and (Plastic or aluminum) (mandatory) Complying with ISO Standard 3826. - Certifications should be supplied by manufacturer's company FDA or CE MARKED . - Plastic film: Medical grad PVC (class VI) complying with ISO specification 3826 (should be supplied by manufacturer's company) - Anticoagulant(mandatory): CPDA1 63ml. - Validity(mandatory) 24 month or more. - Indications:(mandatory) For collection of whole blood, preparation and storage plasma, red blood cell and platelet"&	66823	7.43	اوربي

١. على مقدمي العطاء المؤهلين والراغبين في الحصول على معلومات اضافية الاتصال بوزارة الصحة / الشركة العامة لتسويق الادوية والمستلزمات الطبية(كيمادیا) /قسم الاعلام الدوائي والعلاقات العامة- الطابق الخامس مقر وزارة الصحة البريد الالكتروني dg@kimadia.gov.iq والموقع الالكتروني لكيمادیا www.kimadia.gov.iq والاطلاع على وثائق المناقصة على العنوان أدناه من الساعة الثامنة والنصف صباحا الى الثانية والنصف ظهرا بتوقيت بغداد

٢. **متطلبات التاهيل المطلوبة** :المتطلبات القانونية والفنية والمالية وكما مذكور في وثائق المناقصة.

٣. بامكان مقدمي العطاء المهتمين شراء وثائق المناقصة بعد تقديم طلب تحريري من قبل المدير المفوض للشركة او من يخوله قانونيا (عند او قبل (٢:٣٠ بعد الظهر (٢٠٢٦/٦/٢٢)) وبعد دفع قيمة البيع للوثائق القسم المالي / الطابق السادس غير القابل للرد بمبلغ مقطوع وكالتالي:

أ- مليون دينار عراقي عن المناقصة ذات قيمة مليون دولار او اقل غير قابل للرد.

ب- (٢)مليون دينار عراقي عن المناقصة التي تزيد كلفتها على مليون دولار غير قابل للرد.

ت- ١٥٠ دولار عن المناقصة العامة غير قابلة للرد .

ث- ١٠٠٠ الف دولار عن الاحالات الخاصة بالمستلزمات الطبية (عن طريق الدعوة المباشرة او اسلوب اعطاء الوحيد او الاساليب الاخرى المستثناة من الزام الشركات بتقديم وصل شراء بموجب التعليمات النافذة) .

وبخلافه فان العروض سوف تهمل.

-يعاد ثمن شراء وثائق المناقصات الى المناقصين في الحالات التالية :

أ- عند الغاء المناقصة قبل توقيع العقد ولاسباب مبررة دون تعويض مقدمي العطاءات .

ب- حالة الغاء المناقصة وتغيير اسلوب التنفيذ الى الدعوة المباشرة او العطاء الوحيد .

ت- عند الغاء المناقصات للسنة السابقة والاعلان عنها مجددا وبتسلسل جديد للعام اللاحق .

-العروض التي تصل من خلال البريد السريع فأن المجهز مطالب بتقديم المبلغ المذكور

- (على مقدم العطاءالذي سبق له الاشتراك في المناقصة المعاد اعلانها ان يقدم وصل الشراء السابق لها مع وثائق العطاء). وفي حال تعديل اسعار شراء هذه الوثائق فيتحمل مقدم العطاء الفرق بين السعرين في حال زيادة السعر ويرافق مع عطاءه الوصلين الاول والثاني

٤. يتم تسليم العطاءات على العنوان الاتي (وزارة الصحة / الشركة العامة لتسويق الادوية والمستلزمات الطبية (كيماديا) / الطابق السابع / قسم استيراد المستلزمات الطبية / ادارة صندوق العطاءات / باب المعظم / بغداد / العراق الهاتف :٤١٥٧٦٦٧ , رقم هاتف النقال :٠٧٧٠٥٤١٩٠٧٤ , هاتف البدالة ٨ , ٧ , ٥ , ٤١٥٨٤٠١ , بدالة ذات اربعة خطوط عند او قبل (تاريخ الغلق) [٢٠٢٦/٦/٢٢] لغاية الساعة (٢:٣٠ ظهرا) بتوقيت بغداد وبخلافه لايتم استلام العطاءات المتاخرة وسيتم فتح العطاءات بحضور مقدمي العطاءات او ممثليهم الراغبين بالحضور على العنوان الطابق السادس (لجنة فتح العطائات) في [٦/٢٣ /٢٠٢٦] في الساعة (٩:٠٠ صباحا) بتوقيت بغداد .

٥-على مقدمي العطاءات تقديم التأمينات الاولية المطلوبة مبلغ مقطوع لا تقل عن (١٪) واحد من المائة من الكلفة التخمينية للمناقصة بالدينار العراقي او ما يعادلها بعملة قابلة للتحويل من ضمن قائمة العملات التي يصدر البنك المركزي العراقي اسعار صرفها الى الدينار العراقي

٦-بأمكان مقدمي العطاءات الراغبين في شراء وثائق المناقصة مراجعة جهة التعاقد لغرض شراؤها ورقيا وكما يحق لهم شراؤها (وثائق المناقصة) الكترونيا عبر المنصة الالكترونية الموحدة للاعلانات والمناقصات **IRAQ TENDER PLATFORM** والعائدة لوزارة التخطيط-

- جهة التعاقد غير ملزمة بقبول اوطأ العطاءات
- على مقدم العطاء تقديم العطاءات بنسختين متطابقتين (اصلي ونسخة طبق الاصل) ويثبت نوع النسخة بشكل واضح على شكل مغلف اضافة الى تثبيت رقم وتاريخ المناقصة على كل مغلف

٧- جهة التعاقد غير ملزمة بقبول جميع الكميات في حال اذا كان العرض المقدم من المناقص لجميع الكميات .

٨- يتحمل من ترسو عليه المناقصة اجور النشر والاعلان لاخر اعلان في الصحف الوطنية والمنصة الالكترونية الموحدة للاعلانات والمناقصات واجور توثيق وارشفة العقد الكترونيا غير قابلة للرد (عبر الدخول الى المنصة من خلال الموقع www.itp.iq).

٩- يتم نشر كتاب الاحالة في المنصة الالكترونية الموحدة للاعلانات والمناقصات ويعتبر هذا الموقع مكان للاعلام والتبليغ الرسمي للمناقص الفائز لكافة مقدمي العطاءات المشتركين في المناقصة وتسري مدة الاعتراض على قرارات الاحالة من اليوم التالي للنشر الموقع اعلاه ,

١٠-لمقدم العطاء تحديد مبلغة بشكل نسبة مؤية من الكلفة التخمينية ويعتمد في ذلك جدول الكميات المعد من قبل جهة التعاقد وحب النسبة المحددة وينطبق ذلك على الكلفة التخمينية المعلنة

١١- يكون تاريخ انعقاد المؤتمر الخاص بالاجابة على استفسارات المشاركين في المناقصة في يوم ٦/١٥ /٢٠٢٦

١٢-يلتزم مقدمو العطاءات بمعايير التأهيل والمفاضلة المنصوص عليها في الوثيقة القياسية

١٣-على مقدمي العطاء الالتزام بما تتطلبه الوثيقة بكافة اقسامها

١٤- الموقع على الانترنيت www. Kimadia.gov.iq
dg@ Kimadia.gov.iq

الصيدلاني/
مدير عام الشركة العامة لتسويق الادوية والمستلزمات الطبية (كيماديا)