



EUROPEAN UNION

DELEGATION TO THE REPUBLIC OF SERBIA

Belgrade, 16/07/2010

AG/AP D(2010) 3350

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

Supply of Rabies Vaccines and Equipment for Surveillance of Animal Diseases

Publication ref.: EuropeAid/129809/C/SUP/RS

Tender no08SER01/33/21 + 08SER01/33/31

No	Question	Answer
1.	<p>Lot n°2 : Surveillance equipment and diagnostic kits for rabies</p> <p>2.3 Centrifuge</p> <p>2.3.1. Universal refrigerated centrifuge with rotor</p> <p>1. Is for tubes of 1,5 – 2,2ml necessary angle or swing-out rotor?</p> <p>2. What should be the capacity on rotor for tubes of 1,5 – 2,2ml?</p> <p>3. Which rotor is necessary for tubes of 15 and 50ml – swing out or angle?</p> <p>4. What capacity on rotor is necessary for tubes of 15 and 50ml?</p> <p>2.3.2. Laboratory centrifuge with rotor</p> <p>What should be the capacity on rotor?</p>	<p>Angle rotor</p> <p>Capacity on rotor should be for minimum 24 1.5/2.2 ml tubes</p> <p>Swing-out rotor with adapter for 15 and 50 ml tubes</p> <p>Maximum capacity is 4x250ml</p> <p>Capacity on rotor should be for minimum 24 1.5/2.2 ml tubes</p>
2.	<p>In the procurement notice it is stated that any country that is subject to IPA directive is eligible to the participation.</p> <p>Does it mean that a private entity form Turkish Republic is eligible to participate alone or within a consortium?</p>	<p>Yes.</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
3.	<p>Lot no 3: Vehicles</p> <p>There are EBP and TCF terms in the technical specifications of the vehicle. We could not understand the meaning of these terms as these are not most widely used terms in automotive world.</p> <p>Could you please clarify and give some information for these terms in order to avoid confusion?</p>	<p>Please disregard the reference to EBP and TCF in Technical Specifications for Lot 3.</p> <p>Please note that there will be a Corrigendum No. 2 issued to modify the Technical Specification accordingly and further extend the deadline for submission of tenders for Lot 3 Vehicles.</p>
4.	<p>Lot no.2 Surveillance equipment and diagnostic kits for rabies</p> <p>1. In view of the fact the origin of products must be from EU countries or the countries covered by IPA project, do the European distributors of particular producers outside ED come into account (nasco bags through Deltalab)</p> <p>2. Ordinal number 2.18.2 - in the technical characteristics of ELISA kits for rabies it is not explicitly written whether it is 2 or 5 plates and /or the number of testing per kit.</p> <p>3. Ordinal number 2.13.1-3 the product clarification?</p> <p>4. In Section 7.1 it is written that: The bidder can submit the bid for one lot, several or all lots. Our question is: Must all the items within one lot be offered, and/or will the bids which do not contain all the items within the given lot be correct as well?</p> <p>5. Ordinal number 2.20 -Disinfectant-It is written quantity 12, but unit measure is not</p>	<p>Please refer to the rules of participation and origin which are set out in Article 8 of the Procurement Notice and Sections 3 and 4 of the Instructions to Tender. Tenders must comply fully with both the rules of origin and the rules of participation.</p> <p>Both 2 and 5 strip plates are acceptable.</p> <p>This is a precision cooled rotary saw intended primarily for fine sectioning of a wide variety of samples eg. biomedical samples. It utilizes a replaceable (and fragile) diamond-impregnated blade to cut plastic and bone samples. A variety of sample chucks can hold regularly and irregularly shaped specimens. Liquid fluid or other is using as a coolant/lubricating fluid.</p> <p>All the items within a lot must be offered</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	denoted. Please specify which unit measure is in question.	12 packages for 5 kg each.
5.	Is it admissible the participation of one legal entity in various consortia if they participate for different positions and in this case, is it considered that there is a conflict of interest under section 23.3. from the Instructions for tenderers?	No. A legal entity is not permitted to participate in different consortia for the same contract. Please refer to point number 8 in Section 7 Tenderer's Declaration of the Tender Form for a Supply Contract. The Tender Form for a Supply Contract requires tenderers to declare that they are not tendering for the contract in more than one form. A legal entity may, however, submit a tender for one lot, several or all of the lots.
6.	<p>Lot n 2 Surveillance equipment and diagnostic kits for rabies</p> <p>Question connected with parcel 1.4. on page number 40 in part of tender documentation ANNEX II+III: TECHNICAL SPECIFICATIONS +TECHNICAL OFFER.</p> <p>According your sentence: " The language of the training and the training documents will be Serbian; if not, interpretation/translation will be provided and paid by the Contractor"</p> <p>You asking original manuals (where is that necessary-apparatus, instruments, machines) and they are on English language.</p> <p>Now, please indicate clearly: Is it enough to have original manuals (on English language) and person who will make training can do that orally on Serbian language? If not, please write clearly what exactly you meant.</p>	<p>It is required to supply the original manufacturer's user manuals as well as their translation into Serbian language.</p> <p>The training should be performed in Serbian language. Simultaneous translation of training to Serbian will be accepted. The supplier shall bear all the costs of the training including those for the translation.</p>
7.	<p>Lot 2 Surveillance equipment and diagnostic kits for rabies</p> <p>We didn't find nowhere in tender documentation clearly indicate and because of that we have to ask you next:</p>	

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>1. Does producer (as company), which products we going to offer on this tender bid, must to have ISO Certificate and which ISO Certificate exactly (9001, 14000...)?</p> <p>2. Does all offered products from our side, according request from tender documentation, must to have CE mark?</p> <p>3. If producer (as company), which products we going to offer on this tender bid, must to have ISO Certificate when we have to deliver that certificate ? Is it acceptable to be delivered to Advertiser as COPY (always is just one ORIGINAL and that is for company) and because of so many documentation is possible to deliver that document in the moment of installation and training (If is training provided for that product)?</p> <p>4. If all offered products from our side, according request from tender documentation, must to have CE mark, when we have to deliver that certificate about CE mark? Is it acceptable to be delivered to Advertiser as COPY and because of so many documentation is possible to deliver that document in the moment of installation and training (if is training provided for that product)?</p>	<p>1. Supplier will have to provide certified copies of producer's ISO 9001 and ISO EN 12469 certificates for item 2.4.1 (Biohazard cabinet Class II) as annex to the offer. Producer ISO certification for other items is not required, but in case that producers of other items are ISO 9001 certified, copies of those certificates should also be annexed to the offer.</p> <p>Please note that there will be a Corrigendum No. 2 issued to modify the Technical Specification accordingly and further extend the deadline for submission of tenders for Lot 2.</p> <p>2. All offered products /devices/ from the Tender documentation must have CE mark.</p> <p>Please however note that the supplies must include all necessary parts and accessories (e.g. tubing, cabling, special tools and lab-ware required for regular maintenance and operation) to provide for installation and putting into operation so that the equipment is capable of operation in accordance with these technical specifications and the manufacturers' specifications.</p> <p>All items and sub-items supplied must comply with all applicable Serbian and EU safety standards. Responsibility for obtaining any certificates (e.g. any certificates required from the Serbian Institute of Standardisation) required for the import and putting into use of equipment supplied rests wholly with the Contractor.</p> <p>All electrical equipment must comply with Serbian electrical standards (specifically including power supplies, sockets and plugs). All cabling required for installation and use of the equipment must be included. It is the Contractors responsibility to ensure that all equipment is in fully compatible with the Serbian electrical system.</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
		<p>3. Please see answer to sub-question No. 1 above.</p> <p>4. Certified copies of the CE certification may be requested at the stage of provisional acceptance. Contractors should have them prepared for inspection timely and in advance. Please also see the answer to sub-question No. 2.</p> <p>Please see, previous answers in this section</p>
8.	<p>Lot 2 Surveillance equipment and diagnostic kits for rabies</p> <p>For Part 2, item 2.3.1:</p> <p>1. Is it possible to offer a centrifuge with a maximum capacity: 4x100 ml instead of 4x250 ml?</p> <p>2. Is it possible to offer a complete rotor with 4 places for the tubes of 50 ml and 28 places for the tubes of 15 ml?</p> <p>3. Which is the capacity of the rotor for tubes 1.5 to 2.2 ml which may be offered?</p> <p>For Part 2, item 2.3.2:</p> <p>1. Which is the capacity of the rotor for tubes 1.5 to 2.2 ml which may be offered?</p> <p>For Part 2, item 2.7:</p> <p>1. Is it possible to offer a vortex with a adjustable speed 400-3500 rpm?</p> <p>For Part 2, Item 2.11:</p> <p>1) Is it possible to offer a pipettors (pipette aid) with Cadmium-free NiMH battery powered, direct charging via charger?</p>	<p>NO</p> <p>NO</p> <p>Minimum for 24 1.5/2.2ml tubes</p> <p>Minimum for 24 1.5/2.2ml tubes</p> <p>No</p> <p>No</p>
9.	<p>In the technical specifications requested vehicle is Pick-up car, but it is not clear the body type</p>	<p>Either minivan or cargo van type vehicles that satisfy the technical specifications for Lot 3 will</p>

CONTRACTING AUTHORITY’S CLARIFICATIONS No. 1

No	Question	Answer
	<p>of this vehicle as minivan, cargo van type vehicles can be called as pick up.</p> <p>So we kindly request you to give more information about body type of this vehicle and please confirm if minivan type vehicles are acceptable or not?</p>	<p>be considered without preference.</p>
<p>10.</p>	<p>Lot 2: Surveillance equipment and diagnostic kits for rabies</p> <p>ELISA equipment (2.1.1)</p> <p>Is it acceptable to offer a ELISA washer with linear shaker instead of orbital?</p> <p>What is the minimum number of different buffers acceptable?</p> <p>Microscopes (2.2.1)</p> <p>In the specifications it states that the eyepiece needed should be “Pair of 10x – FOV18mm”. Is it acceptable to offer eyepiece 10x magnification with larger field of view (FOV)?</p> <p>In the specification it states that the achromatic objective needed should be 63X. Is it acceptable to offer a more common objective with magnification of 60X?</p> <p>Laminar flow (2.4.1 and 2.4.2)</p> <p>Is it acceptable to offer a laminar air flow cabinet with depth of 86cm?</p> <p>Is it acceptable to offer a laminar air flow cabinet with depth of 86cm?</p> <p>Incubator (2.5.1)</p> <p>If the incubator meets the temperature range and</p>	<p>Orbital shaker is required.</p> <p>Minimum 3</p> <p>YES</p> <p>NO</p> <p>NO</p> <p>NO</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>stability specifications, is it possible to offer an incubator with 1400 Watt heater?</p> <p>Freezers (2.6.3)</p> <p>In the specification it states that the capacity of the refrigerator must be at least 1300L. Is it acceptable to offer 1240L capacity?</p> <p>Pipettors (2.11.1)</p> <p>Is it acceptable to offer NiMH battery instead of NiCd batteries?</p> <p>Electrophoresis (2.8.1)</p> <p>Is it acceptable to offer a electrophoresis unit with different size of the gel for minimum 80 samples (size being 14.4 x 10.2 cm)?</p> <p>Does the electrophoresis unit offered need to include power supply?</p>	<p>1500 Watt heater is required.</p> <p>NO</p> <p>NO</p> <p>NO. Required gel dimensions are specified in Technical Specifications for Lot 2.</p> <p>NO. Please refer to Technical Specifications for Lot 2.</p>
11.	<p>Lot n° 2: Surveillance equipment and diagnostic kits for rabies</p> <p><u>Question 1:</u></p> <p><u>For 2.13 (2.13.1 2.13.2 2.13.3)</u></p> <p>Regarding of our market research this Low Speed Saw Unit in the technical specifications is not a product that originated of European Unions countries so could you please give derogation for this product?</p>	<p>NO</p>
12.	<p>LOT 3 – Vehicles</p> <p>1. In the technical specification there are abbreviations EBP and TCF. Can you please clarify to us the meaning of those two abbreviations, given that different car manufacturers do not have the same terminology in terms of characteristics of the vehicles.</p>	<p>Please see Answer n° 3.</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>2. In the technical specification is given that motor should be 1.4. Does this mean that engine volume must be greater than 1400 cm³ or that label provided by the manufacturer must be 1.4 or higher?</p> <p>3. Usually manufacture cycle is 60 days for standard cars with standards equipment. These 12 cars should be equipped with cooling department (absolute non standard request) which extends manufacture for 10 days per car. Is it possible to get extension of delivery period for this LOT up to min 120 days?</p>	<p>Minimum engine volume must have manufacturer's label 1.4 (engine volume of 1400 cm³ +/- 50 cm³) or higher.</p> <p>The implementation period for Lot n° 3 shall be within 120 days after the Commencement order.</p> <p>Article 18 of General Conditions of Contract defines the Commencement order.</p> <p>Please note that there will be a Corrigendum No. 2 issued to the tender dossier to modify the implementation period and further extend the deadline for submission of tenders for Lot 3 Vehicles.</p>
13.	<p>Technical Specifications – 3.1</p> <p>- It is required that a vehicle has ‘EBP’ and ‘TCF’. As there is more than one explanation of these abbreviations, would you be so kind to state exact meaning of them, i.e. explain what they stand for?</p> <p>- It is required that there is a ‘air-conditioning installed’. Is this requirement for the driver and passenger cabin or for the load compartment?</p> <p>Special conditions</p> <p>- Article 13 and 19 – Programme (period) of implementation of the tasks: it is stated that ‘the implementation period, ending with Provisional Acceptance, shall be, for Lot 3: within 2 months after the Contractor receives the Commencement order’.</p> <p>Please clarify when is the Commencement order to be given / received – is it the date of the Contract signature or some later date, depending on fulfillment of some other requests?</p>	<p>Please see Answer n° 3.</p> <p>Condition “Air-conditioning installed” refers to driver / passenger cabin. Load compartment (closed and isolated cooling compartment) has its particular conditions as described in technical specifications, i.e. cooling compartment should enable temperatures of 4 C (+/-1.5 C).</p> <p>Please see answer to clarification request n° 12.</p>
14.	<p>1. Lot 3, item 3.1 – pick-up car with cooling compartment installed – the technical specifications requests EBP and TCF. Please advise what these acronyms stand for.</p> <p>2. Lot 2, item 2.16.2 – bag for transport of sample – please indicate size/volume of the bags</p>	<p>Please see Answer n° 3.</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>required.</p> <p>3. Lot 2, item 2.17 – reagents 2.17.1 – 2.17. 4 – please advise required volume of each reagent</p> <p>4. Lot 2, item 2.18 – Diagnostics – items 2.18.1 – 2.18.2 – the specifications included in the tender dossier is insufficient, please provide us with further details.</p> <p>5. Lot 2, item 2.20 – Disinfectant – please advise required volume of disinfectant.</p>	<p>Min. 22x15 cm</p> <p>2.17.1 40 bottles a 500ml</p> <p>2.17.2 4 bottles a 500 ml</p> <p>2.17.3 22 bottles a 100 ml</p> <p>2.17.4 3 bottles a 100 ml</p> <p>2.18.1 70 packages</p> <p>2.18.2 12 kits - 2 plates for 192 reaction per kit</p> <p>12 packages a 5 kg</p>
15.	<p>Lot n° 2: Surveillance equipment and diagnostic kits for rabies</p> <p><u>Question 1:</u></p> <p><u>For 2.2.1 Inverted Fluorescence Microscope</u></p> <p>It is requested that the system should have an ABBE type condenser with a N.A value of 1.25. According to our 20 years of experience on microscopy, I would surely say that none of the Inverted microscopes are equipped with Abbe type 1.25 condensers. This type of condensers is mainly used in “Upright” microscopes.</p> <p>Would you please clarify this situation for us in order to provide an appropriate offer?</p>	<p>Condenser should be LD N.A 0.55H, Ph1, Ph2, Ph3, DIC, DIC</p> <p>Please note that there will be a Corrigendum No. 2 issued to modify the Technical Specification accordingly and further extend the deadline for submission of tenders for Lot 2.</p>
16.	<p>Lot n 2 Surveillance equipment and diagnostic kits for rabies</p> <p>1. Could you explain us more about reference table which is included in tender form for supply contract? Precisely, we wonder whether is it enough to fill that table just for one client or</p>	<p>1. It is enough to complete the table in the prescribed format which serves to summarise the major relevant supplies carried out in the course of the past 3 years by the legal entity or entities making the tender. Please be aware that the</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>to copy that table for representing more clients in order you could see our complete reference?</p> <p>2. How do we fill Tender form for supply contract, point 7. Tenderer(s) Declarations? Do we need to make separate declarations for every point (1-14, especially for point 2. and 3.) or we have to include all of these statements within a given form?</p> <p>3. In Instructions for tenderers point 11, part 3 "Documentation" you stated that we need to submit duly authorised signature: an official document (statutes, power of attorney, notary statement, etc.) proving that the person who signs on behalf of the company is duly authorised to do so. Our question: is OP form (<i>OP obrazac</i>) sufficient or we have to submit OP form, together with our statute?</p> <p>4. Do we have to include in dossier following documents (which are included in tender dossier, but point 11. of Instructions for tenderers doesn't mention them): evaluation grid, administrative compliance grid, General and Special conditions, delivery list, acceptance?</p> <p>5. Do we have to send pre-financing guarantee and performance guarantee in our offer?</p> <p>6. Could you clarify term "commercial warranty" because we don't have that term in our law, is it different from warranty according to Serbian law? Especially according to terms in point 14. and 15. in detailed administrative compliance grid, what are the difference between these two warranties?</p> <p>A description of the organisation of the warranty tendered, which must be in accordance with the conditions laid down in Article 32 of the General Conditions</p> <p>A description of the organisation of the commercial warranty tendered in accordance with the conditions laid down in Article 32 of the Special Conditions</p> <p>7. Could you explain us, please, what kind of</p>	<p>table can include references from more than one client. Please note that the total number of references provided must not exceed 15.</p> <p>2. As part of their tender, each legal entity identified in Section 1 of the Tender Form for a Supply Contract, including every consortium member, must submit a completed and signed declaration in the exact format provided in Section 7 of the Tender Form for a Supply Contract. All of the points/statements must be included in the tenderer's declaration. Separate declaration forms for every point are not required. Please however note that one completed and signed Tender Form for a Supply Contract must be supplied for each lot if the legal entity is tendering for more than one lot.</p> <p>3. Whatever official document is submitted, whether it is the statutes, a power of attorney, a notary statement, the OP form (<i>OP obrazac</i>) etc, it must prove that the person who signs on behalf of the company/joint venture/consortium is duly authorised to do so.</p> <p>4. No.</p> <p>5. No.</p> <p>6. The description of the organisation of the warranty required by the Contracting Authority, i.e. the minimum warranty conditions, is set out in Article 32 of the General Conditions. Article 32 of the Special Conditions further amplifies and supplements the minimum warranty requirements of the Contracting Authority set out in Article 32 of the General Conditions. The complete list of conditions and the period of the warranty, namely 24 months, required by the Contracting Authority for Lot 2 and Lot 3, is thus described in the abovementioned articles of the General and Special Conditions. Therefore the warranty offered by tenderers must satisfy all warranty requirements defined in Article 32 of the General Conditions and Article 32 of the Special Conditions.</p> <p>7. The kind of proofs required to prove that tenderers are not in any of the exclusion situations which would exclude them from</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p>proofs are those mention in Tender form for supply contract, point 7. Tenderers declaration(s), precisely declaration nr. 9 (proofs according to section 2.3.3 of the Practical Guide to contract procedure for EC external actions and proofs according to section 2.4.11.1.3 and 2.4.11.1.4 of the Practical Guide)?</p> <p>We will not send that proofs, as far as we understand it, in our offer, but only in the event of successful offer (after notification)?</p> <p>Is statement in point 13. of Detailed administrative compliance grid (a statement to the effect that tenderer is not in any of the situations listed in section 2.3.3 of the Practical Guide to contracts procedures EC external actions) refers to declaration nr.9, point 7. in Tender form for supply contract ?</p> <p>8. How do we prove "Eligible origin and proper statement attesting the origin of the supplies tendered" (point 3. in Detailed administrative compliance grid)? Do we submit statement in form Annex V "Statement of origin" which is included in tender dossier, or we have to prove that in different form, especially according to point 11. Instructions for tenderer(s) where is stated that we need to supply on free format a statement by the tenderer attesting the origin of the supplies tendered (or other proofs of origin)?</p> <p>9. Do we submit a project implementation schedule with major milestones in a free format? (point 16. Detailed administrative compliance grid).</p> <p>10. In relation to product: 2.18.1 Flourescin (FITC) conjugate anti-Rabies For Rabies Immuno diagnosis using fluorescence test</p> <p>Is it acceptable to offer this product in packages of 1 ml (antirabic conjugate) + 1 ml (negative)? Accordingly, is volume 1+1 ml acceptable?</p>	<p>participating in these contracts are listed in Section 2.3.3 Grounds for Exclusion of the Practical Guide to Contract Procedure for EU External Actions. Please therefore see Section 2.3.3 of the Practical Guide to Contract Procedure for EU External Actions.</p> <p>The kind of documentary proofs required to prove the tenderer's financial and economic standing and the technical and professional capacity according to the selection criteria for this call for tender are listed in Sections 2.4.11.1.3 and 2.4.11.1.4 of the Practical Guide to Contract Procedure for EU External Actions.</p> <p>The abovementioned documentary proofs must be submitted after notification of the contract award.</p> <p>Yes.</p> <p>8. Yes. For the proof of eligible origin and proper statement attesting the origin of the supplies tendered tenderers must submit a statement in the format given in Annex V Statement of Origin.</p> <p>9. Yes.</p> <p>10. That would be one of the possibilities.</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
17.	<p>Lot 2: Surveillance equipment and diagnostic kits for rabies</p> <p><u>Lead rubber apron (2.15.1)</u> What is the dimensions of the apron needed?</p> <p><u>Bag for transport of sample (2.16.2)</u> What is the volume of the bag needed?</p> <p><u>Dulbecco modified medium (2.17.1)</u> Can you please specify in more detail precise concentrations of the medium as well as the packaging volume?</p> <p><u>Foetal bovine serum (2.17.2)</u> Can you please specify packaging volume needed?</p> <p><u>Trypsine (2.17.3)</u> Can you please specify packaging volume needed?</p> <p><u>Antibiotic / Antimycotic solution for cell culture (2.17.4)</u> Can you please specify packaging volume needed?</p> <p><u>RNA extraction kit (2.17.5)</u> Can you please specify number of reactions in the single packaging needed?</p> <p><u>OneStep RT-PCR kit (2.17.6)</u> Can you please specify which cycler is used and if the detection is made by probes or by SYBER green? Can you please specify number of reactions in the single packaging needed?</p> <p><u>PCR purification kit (2.17.7)</u> Can you please specify number of reactions in the single packaging needed?</p> <p><u>Flourescin (FITC) ant-rabies conjugate</u></p>	<p><i>L and XL</i></p> <p>Min. 22x15 cm</p> <p>1X ready to use mix a 500ml</p> <p>a 500ml</p> <p>a 100 ml</p> <p>a 100 ml</p> <p>minimum 250 reactions/kit</p> <p>It is kit for Reverse Transcriptase PCR not for real time PCR</p> <p>minimum 100 reactions/kit</p> <p>minimum 250 reactions/kit</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	<p><u>(2.18.1)</u> Can you please specify number of reactions in the single packaging needed?</p> <p><u>Microscope slides (2.19.1)</u> Can you please specify quantity of microscope slides needed in a single packaging?</p> <p><u>Cover glass (2.19.2)</u> Can you please specify quantity of cover glass slides needed in a single packaging?</p> <p><u>Disinfectant (2.20)</u> Can you please specify volume of disinfectant needed in a single packaging?</p>	<p>At least 120 reactions/packaging(unit)</p> <p>50 per pckg/box</p> <p>100 per pack/box</p> <p>12 packages a 5 kg</p>
18.	<p>Lot n 2 Surveillance equipment and diagnostic kits for rabies</p> <p>For Part 2, item 2.2.1:</p> <p>1. Do You mean upright microscopes? Inverted microscopes do not have Condenser: Abbe, N.A. 1.25.</p> <p>For Part 2, item 2.3.1:</p> <p>1. Is it possible to offer a pipettors (pipette aid) for Pipetes types - glass and plastic pipettes from 1-100 ml with 0,2 filters.</p> <p>2. Is it possible to offer a pipettors (pipette aid) with Cadmium-free NiMH battery powered, direct charging via charger?</p> <p>For Part 2, item 2.6.2:</p> <p>1. Is it possible to offer incubator with alarm system: monitor connected to a computer to pull out the information with own power source which will work if the cabinet is without power and with possibility to connect it for tex message alerts.</p> <p>For Part 2, item 2.12.11:</p> <p>1. Is it necessary and is it obligatory to offer</p>	<p>Condenser should be LD N.A 0.55H, Ph1, Ph2, Ph3, DIC, DIC</p> <p>No (the question seems to relate to item 2.11).</p> <p>No (the question seems to relate to item 2.11).</p> <p>YES</p> <p>NO</p>

CONTRACTING AUTHORITY'S CLARIFICATIONS No. 1

No	Question	Answer
	sealing foil for PCR plates?	