REQUEST FOR TECHNICAL ADVICE MEETING FOR CERTIFICATION OF SUITABILITY

to be filled in for each request for a Technical Advice meeting related to the procedure for Certificate of Suitability to the monographs of the European Pharmacopoeia AP-CSP (07) 1

1	General information	
1.1	Date of submission of the request for Technical Advice meeting:	dd/mm/yyyy
1.2	Dates proposed for the technical advice meeting (at least 2, preferably 3):	
	This form should be submitted to EDQM one month before any proposed meeting dates.	
1.3	Organisational matter - tick box, as appropriate:	
	Meeting at EDQM premises or Teleconference	
2	Scope for the questions for technical advice - <i>tick box(es), as a</i>	ppropriate.
	Implementation of resolution, procedures and policies for certific	ation
	Application(s) for new certificate(s) of suitability	
	Revision(s) or renewal(s) of existing certificate(s)	
	CEP number(s): Name(s) of substance(s) + subtitle (if applicable):	
	Preparation of answers to deficiency letter	
	CEP number(s):	
	Name(s) of substance(s) + subtitle (if applicable):	
	Other, please specify:	
_	tions and relevant documentation shall be attached; the reques ng (see annex 2)	t cannot be accepted if
The	relevance of the Technical Advice request will be decided at rec	ceipt by the EDQM.

3	Names and addresses					
3.1	Contact person for this request for technical advice					
	Title and surname:		First	name:		
	Job title:		Department:			
	Tel:		Fax:			
	E-mail:					
	Name of the company applying for the Technical Advice:					
	Address:					
	Postcode:		Town:			
	Country:					
	Tick this box if the company above is different from the (intended) CEP Holder and include annex 1					
3.2				meeting (Max 4). If differ the his/her presence	erent from the company	
	above mentioned please specify the link and justify his/her presence. If not yet available that should be completed and confirmed at the latest 10 days before the meeting.					
	Title and surname	First name		Name of the company	Department	

4. This form and questions should be sent in electronic format to: CEP@edqm.eu

European Directorate for the Quality of Medicines & HealthCare Certification of Substances Department

5. Invoicing details (mandatory)

Following receipt of the request for technical advice EDQM will send you an invoice. Please proceed with payment <u>after</u> you receive the invoice.

Reference	Item	Price
CEP11	Technical advice	1000€

Contact person, authorised for communication on behalf of the company:		
Title* (Mrs, Mr, Dr)		
Contact first name*		
Contact family name*		
Job title/Department		
Company name*		
Address*		
Postcode*		
City*		
Country*		
Telephone*		
Fax*		
Email*		

Fields marked * are mandatory

	INVOICING ADDRESS
COMPANY DETAILS	
EDOM Client Code	
Company name (*):	
Address(*)	
City (*):	
Postcode (*)	
Region/State	
Country (*)	
VAT Number (**)	
Tel (switchboard) (*)	
Fax (*)	
Email (*)	
Contact name(*)	
Contact first name (*)	

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Job title (*)	
Department (*)	
Tel (*)	
Fax (*)	
Email (*)	
Fields marked (*) are mandatory. Field	ds marked with (**) are required for EU only.
PREFERRED LANGUAGE (for inv	voicing/accounting only):
☐ English	
☐ French	
<u>PAYMENT</u>	
	al advice meeting, we will send you an invoice. Please note n 30 days end of month. Details of payment methods will be able to settle your invoice by:
1. BANK TRANSFER 2. CREDIT CARD	

European Directorate for the Quality of Medicines & HealthCare Certification of Substances Department

Annex 1

Template for letter of Authorisation

[address of the (intended) CEP Holder]

[date and place]

LETTER OF AUTHORISATION

We, [name of the (intended) CEP Holder], hereby authorise, [name of the authorised representative], to act as contact for our request for technical advice for Certificate(s) of Suitability for [name(s) of the substance(s)].

Signature

Annex 2

Technical Advice questions & documentation: (To be filled in if not, the request cannot be accepted. Full supportive documentation should also be provided, if needed, as annexes)		