



Office of the Deputy Director-General S/248/2001 19 March 2001 ENGLISH only

OPCW SURVEY ON THE CAPACITIES OF MEMBER STATES REGARDING THE HANDLING AND ANALYSIS OF BIOMEDICAL SAMPLES

Background

- 1. The Chemical Weapons Convention provides for the collection of samples of biomedical origin during investigations of alleged use of chemical weapons by the OPCW (Verification Annex, Part XI, paragraphs 16 and 17). Analysis of these samples may provide information that contributes to the inspection team's conclusions regarding allegations of the use of chemical weapons. The list of approved equipment adopted by the Conference of the States Parties includes an alleged use sample collection kit, with a module for the collection of biomedical samples (annex, attachment 16, to C-I/DEC.71 and Corr.1, both dated 23 May 1997).
- 2. While the Preparatory Commission and the OPCW both devoted considerable resources to the development of procedures and a network of accredited laboratories for the collection, transport and analysis of environmental samples, similar arrangements are not yet in place for samples of biomedical origin.
- 3. As part of an exercise to assess current international abilities in this field, and in order to assist decisions on development work that may be required, the Secretariat would greatly appreciate receiving answers to the questionnaire annexed to this document. National Authorities are requested to forward copies of this questionnaire to laboratories that they consider may have relevant abilities in the field of analysing samples of biomedical origin for the presence of chemical weapons. Responses should be sent, if possible, before 12 April 2001 to:

Health and Safety Branch OPCW Secretariat Johan de Wittlaan 32 2517 JR The Hague The Netherlands

Annex: Questionnaire on national capacities for the analysis of biomedical samples

Annex

NATIONAL CAPACITIES FOR THE ANALYSIS OF BIOMEDICAL SAMPLES Questionnaire

Laboratory name:						
Address and telephone:						
Contact pers	on:					
	lology for retros vailable in your		n of toxic sch	neduled chemicals in biomedical		
Yes / No						
If yes, ple	ase continue:					
		als (or their meta samples at your l		eakdown products) can currently		
presence note in ea	of scheduled che ch case whether	emicals (or their	metabolites ar	yse biomedical samples for the nd breakdown products)? Please ated or not, in which species, and and tissue.		
Met Example: Acety activit	lcholinesterase	<u>Validated</u> Yes	Species Human	Blood / Urine / Tissue Blood		

4)	Does your laboratory have internationally recognised quality control accreditation for the analytical techniques involved in biomedical sample analysis?
	Yes / No
	Comments:
5)	Is your laboratory currently active in research into techniques for analysis of biomedical samples containing scheduled chemical or their metabolites?
	Yes / No
	If yes, please provide a separate list of relevant published references, and if possible, copies of relevant publications.
6)	Do you have experience regarding the storage and transportation of biomedical samples from persons possibly exposed to scheduled chemicals?
	Yes / No
	If yes, please provide details:
7)	Does your laboratory have experience of handling and analysing biomedical samples originating from actual incidents of suspected chemical weapons use?
	Yes / No
	If yes, please provide details:

8)	Is your laboratory willing to be designated by the Director-General of the OPCW for the analysis of biomedical samples in the context of OPCW activities, and are you willing to participate in inter-laboratory comparison exercises and proficiency testing?
	Yes / No
	Comments:
9)	Is your laboratory willing to share knowledge and skills regarding the analysis of biomedical samples (via, for example, provision of training to Nationals of other Member States)?
	Yes / No
	Comments:
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