

EUROPEAN COMMISSION

HEALTH & CONSUMER PROTECTION DIRECTORATE-GENERAL

HIGH LEVEL GROUP ON HEALTH SERVICES AND MEDICAL CARE

Document: Minutes of the third meeting of the High Level Group on 13 October

2004

To: Members From: Secretariat

Action: For information.

This revised version incorporates comments from delegations on the original

draft.

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HEALTH & CONSUMER PROTECTION DIRECTORATE-GENERAL

Brussels, 18 October 2004 HLG/2004/19 REV1

THIRD MEETING OF THE HIGH LEVEL GROUP ON HEALTH SERVICES AND MEDICAL CARE 13 OCTOBER 2004, BRUSSELS

Subject: Minutes of the meeting

1. WELCOME AND INTRODUCTION

Participants were welcomed to the third meeting of the High Level Group by the chairman, Mr Madelin. A list of participants is attached. The draft agenda was adopted as proposed.

2. SPECIAL PLENARY SESSION

This exchange of views began with a detailed presentation by Mr Schaub, Director-General of DG MARKT, outlining the proposal for a directive on services in the internal market¹ and the aspects of the proposal related to health services.

In discussion, members supported the aims of the proposed directive, but felt that it did not take specific account of the particular nature of health services and the non-market environment in which publicly-funded health services were largely provided. Although there was not sufficient time to go into points of detail, particular issues raised included:

- the application of country of origin rules to health services risked undermining the ability of Member States to maintain their mechanisms for ensuring quality of health services provided on their territory and planning service provision;
- the distinction between hospital and non-hospital care was counterproductive in health policy terms, where the emphasis was increasingly on integrated packages of care. This would only continue with greater numbers of older people and the consequent increased focus on the management of chronic disease, and legal distinctions between hospital and non-hospital care risked distorting care provision away from the most medically appropriate approach;

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¹ See COM(2004)2.

- health systems were complex and delicately balanced structures, and applying simple general rules such as those in the proposed services directive risked upsetting that balance in unforeseeable ways that could undermine the basic values of accessibility, universality and equity on which they were based;
- the organisation and financing of health services being a responsibility of the Member States, the mechanisms of the directive providing the Commission and the Court with a scrutiny role over those services were inappropriate.

It was agreed that these discussions and responses to the issues raised by delegations should be pursued in more detail within the Council, and Mr Schaub emphasised the availability of his colleagues to discuss detailed issues. However, Mr Schaub remained unconvinced that health services were so specific that solutions to the issues raised could not be found within the framework of the services directive, and called on Member States to approach discussions on that basis.

3. MINUTES OF THE LAST MEETING

The draft minutes of the first meeting (HLG/2004/16) were approved, subject to an amendment to reflect the inclusion of Spain in the countries who had welcomed the UK initiative on patient safety and revised text proposed by Belgium to clarify the scope of the working group on cross-border healthcare purchasing and provision.

4. REVIEW OF PROGRESS IN WORKING GROUPS

Cross-border healthcare purchasing and provision

It had been agreed that this working group would be chaired jointly by Malta as well as the Netherlands. The working group had made a roadmap of future work, and would focus on what package of care people were entitled to, the financial impact of cross-border care, legal responsibility and questions around 'undue delay'. As the responses of the Member States to the questionnaires and the descriptions of the experiences in the member states were received shortly before the meeting, the content of the contributions could not be discussed and conclusions could thus not yet be drawn.

The Europe for Patients research project had also been presented to the working group, and could provide useful input. The working group had also considered the German proposal for a conference during 2005; it was felt that the conference should be over more than one day in order to adequately address issues, but this would require additional financial support. The Commission also suggested that the working group should discuss their planned work with the Administrative Commission for Migrant Workers, and draw on their expertise on cross-border care issues.

Health professionals

This working group had met in the morning for the first time, and focused on identifying areas of added value, taking into account the work of other bodies such as the Committee of Senior Officials on Public Health and current work on

exchanging information on poor performing professionals already being taken forward under the Dutch Presidency. To this end, the working group agreed to focus on just doctors and nurses, and to concentrate on three areas:

- better evidence on current practice and current distribution and mobility of health professionals within the Union;
- the impact of professional migration on their health systems of origin and ethical recruitment practices;
- and quality issues, in particular continuing professional development.

It might be useful to commission a study or other work in order to gather better data on these issues; the existing mechanisms for analysing data on professional mobility within the Commission's Internal Market Directorate-General could also help. The European bodies representing doctors and nurses might also be able to provide useful data, and current work by the World Health Organisation (WHO) should also be taken into account.

Centres of reference

Responses to the questionnaire circulated after the first meeting had been received from most members of the working group, showing that there was already a reasonable similarity of view on this topic. Before proceeding to the stage of mapping centres of reference, work was needed to agree on the principles of the definition of a centre of reference, how centres should be designated (at what level, by what process, for what duration) and how they should be organised (networking mechanisms, referral procedures).

It would be important to keep in mind how any European system of centres of reference could add value to existing national systems; data on national systems for centres of reference from other Member States would also be very welcome. There were also major links between the issues being addressed by this working group and several others, such as health professionals, cross-border healthcare, and information and e-health.

The next stage would be to analyse the responses to the questionnaire and to develop a paper setting out areas of consensus and alternative scenarios for areas of difference, taking into account existing models of European cooperation in other areas (such as the Community Reference Laboratories for animal and plant health). This would be discussed at an additional meeting of the working group to be held on 27 October in Brussels.

Health technology assessment

After the first meeting a revised version of the paper setting proposals for networking in this area had been prepared and discussed. A pilot project on networking would aim to make progress in six areas:

 methods for developing common core information packages, including the positive and negative health implications, economic information and socioeconomic and organisational issues to be taken into account within health systems;

- transferability of health technology assessments;
- commissioning reports of common core information on health technology assessment;
- quality management procedures for producing health technology assessments;
- tools for identifying needs and prioritising topics for European health technology assessment;
- and tools for tailoring common core information to inform health policy within Member States.

Developing European-level networking on health technology assessment was not intended to displace national bodies for health technology assessment, but rather to maximise synergies and avoid duplication between them, as well as providing a base of expertise which could be used by Member States to develop their own internal capacities and policies.

Although there was a large measure of consensus on the ideas set out in this paper, it was quite a technical document and thus not necessarily suitable as a report to the Council, and a separate short reporting document was also therefore being prepared. This would need to make clear the added-value of European cooperation on health technology assessment (one example mentioned was that of breast cancer screening, where ten Member States had carried out their own separate technology assessments), whilst recognising that national systems were different and thus evidence and recommendations would have to be adapted to the systems in question.

Information and e-health

This working group was focusing on the issues raised in the recent Commission Communication COM(2004) 356 "E-Health – making healthcare better for European citizens: an action plan for a European e-Health Area" and how best these could be taken forward. A framework of action for addressing these issues had been agreed; one key issue would be funding and how to develop European solutions in this area.

For the time being, this working group was focusing more on the structures and methods of exchanging information through e-health rather than the content of information itself. It would be useful, however, to include a reference to the work that was being undertaken on that area, for example under the health information strand of the public health action programme, which the Commission could provide.

Health impact assessment and health systems

A revised version of the initial paper had been circulated and discussed by this working group; this would be further revised in the light of the morning's discussion. The principles of health impact assessment and health systems were being developed; the next stage would be to propose the development of a framework for assessing impacts of other Community policies on health systems and to identify policy areas where this framework could be applied, in order to

provide information to the European institutions on the impact of proposals on health and health systems.

This was an area which could be very helpful in addressing some of the concerns raised by ministers and other stakeholders in the context of the patient mobility reflection process. However, it would be important to ensure that action was solidly rooted in the Treaty provisions, in particular the requirement of Article 152 that "a high level of human health protection shall be ensured in the definition and implementation of all Community policies and activities", and that the responsibilities of different bodies working in this area were clearly defined.

5. OUTLINE REPORT TO COUNCIL AND NEXT STEPS

The Commission had circulated an outline draft report (HLG/2004/18) including a template for the contributions of working groups; this seemed broadly along the right lines (though it should focus more on output and decisions that ministers should take, rather than activity). The precise institutional mechanism of the report and the Council's response would also need further consideration, but the basis of the report would be that it reflected the views of the members of the High Level Group. To ensure that this was the case, the meeting on 4 November 2004 would be crucial, and a revised and completed draft should be circulated well before that meeting to allow proper consideration.

In terms of follow-up action, members were invited to send any comments on the draft report (HLG/2004/18) by noon on 20 October 2004. Draft contributions from working groups should be provided by Friday 22 October in order to enable a revised and completed draft report to the Council to be circulated in advance of the next meeting of the High Level Group on 4 November.

On the timing of that meeting, health attachés would be in Geneva for negotiations on the International Health Regulations, but it was agreed to proceed with the meeting in any case.

6. OTHER BUSINESS

No items of other business were discussed.

HIGH LEVEL GROUP ON HEALTH SERVICES AND MEDICAL CARE

Subject: Participants in the third meeting of the High Level Group, 13

October 2004

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