EUROPEAN COMMISSION
JOINT RESEARCH CENTRE

Directorate F - Health, Consumers and Reference Materials (Geel)
Health in Society

NCD-Indicators collection process – Diabetes domain Working Group

1st Physical Meeting, JRC Ispra, 4th November 2022

Annotated preliminary agenda

8:30-9:00 Reception

9:00-9:30 Welcome (N.Nicholson)
9:30-9:50 Objectives of the EU NCD Initiative (N.Nicholson)

9:50-10:10 Planning a Scoping Document: Rationale and procedure (F.Carinci)

10:10-11:00 Plenary Discussion

11:00-11:30 Coffee break

11:30-13:30 **Breakout sessions**

Three different breakout sessions will discuss in more details the contents of the three major streams considered for the Scoping document. The objective of these sessions is to provide input for three dedicated chapters included in the document. Participants to the sessions will be pre-assigned by three different facilitators in advance of the meeting. The facilitators will compile specific background materials for the sessions and use their preferred procedure to deliver the final results. Each facilitator will write a detailed report of each session, contributing to the contents of the scoping document.

Stream A. Stakeholders' forum (M.Massi Benedetti)

In this session, participants will identify the criteria relevant to the NCD initiative allowing to classify the major outputs needed by the primary stakeholders. To this end, participants will need to identify classes of users, and assign to each of them a group of outputs e.g. indicators, open data etc. Diabetes will be used as a practical case study in which the different possible outputs will be allocated. The taxonomy will be used to provide input for methods and tools in the final scoping document. The group will also indicate three different outputs that could be tested in a rapid pilot study in diabetes.

Stream B. Methods and tools (F.Carinci)

In this session, participants will identify the main methodological challenges that must be tackled by the information infrastructure. Learning from recent experiences in the EUBIROD network, participants will indicate which developments should be foreseen, what are the limitations of data collection at the data source level (e.g. data linkage, consent, etc), and which data management and statistical tools should be included in the

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future development of a stable framework for the European Union. The identified methods and tools will be at least partly applied for the pilot study.

Stream C. Privacy-by-design (C.T.Di Iorio)

In this session, participants will discuss the latest trends in European legislation regarding the sharing of health data, including potential synergies with the EHDS. The sustainability of the information infrastructure will be discussed, learning from recent experiences from other disease areas and activity at the JRC (e.g. ECIS and ERDRI). The outputs will be used to provide input for other streams in the Scoping Document, including the possible arrangements for the institutionalization of the framework.

13:30-14:30 Lunch break

The wrap-up session will share the results and provide recommendations to the facilitators for the preparation of the Scoping Document.

Wrap-up from Breakout sessions
14:30-14:45 **Who needs what? Stakeholders' forum (M.Massi Benedetti)**
14:45-15:00 **Which information infrastructure? Methods and tools (F.Carinci)**
15:00-15:15 **What is legally viable? Privacy-by-design (C.T.Di Iorio)**
15:15-16:00 Discussion

16:00-16:15 Coffee break

16:15-17:00 Planning a pilot study for diabetes (F.Carinci)

17:00-17:30 Wrap-up (N.Nicholson)