

# Original Clinical Document Evaluation Form for the Functional e2e testing MyHealth@EU Spring 2025 Test Session

Fields marked with \* are mandatory.



**My health @ EU**  
eHealth Digital Service Infrastructure  
A service provided by the European Union

## Evaluation Form for the Functional end-to-end Testing - Original Clinical Document

### Introduction

**The functional end-to-end testing validates, from the user point of view, the process and the information presented to health professionals using the MyHealth@EU services.**

The evaluation is carried out for all the services in an environment that intends to emulate the normal operation as much as possible. For more details on the methodology of the testing, please check the dedicated page [here](#).

The tool used to collect the feedback from the health professionals and semantic experts participating in the testing is the Evaluation Form, which is provided using the European Commission online tool EUSurvey.

Thank you very much for your participation.

### 1 Information on the participants in the evaluation

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\* 1.1 Please, indicate the NCPeH you are representing

IE - Ireland

\* 1.2 Role of the Healthcare Professional

- ☐ Doctor of medicine  
☒ Nurse

- ☐ Dental practitioner
- ☐ Midwife
- ☐ Pharmacist
- ☐ Pharmacist technician
- ☐ Other

\* 1.4 Professional role of the Semantic expert

Technical Architect

## 2 Information on the Evaluation

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\* 2.1 Date of the evaluation

10/04/2025

\* 2.2 Country of origin of the OrCD document being evaluated

LU - Luxembourg

## 3 Identification of the patient process

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### 3.1 Identification of the patient

3.1.1 Could the patient be identified using the fields provided by the country of origin?

(whether the returned information was sufficient to identify the patient)

- ☒ Yes
- ☐ No

3.1.3 Did you encounter any technical error during the Patient Search?

- ☐ Yes
- ☒ No

### 3.2 Identification of Authorized Third Party (Next of Kin)

(optional; Section only applicable when the identification of Authorized Third Party is supported by the implementations in both countries participating in the test)

3.2.1 Is your country implementing the identification of the Authorized Third Party (Next of Kin) operation?

- ☐ Yes
- ☒ No

## 4 Non-Functional Requirements

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\* 4.1 Was the system available during the performance of the test?

- ☒ Yes  
☐ No

#### 4.3 Response Time

	>15s	<15s	<10s
* To find the patient	<input type="radio"/>	<input checked="" type="radio"/>	<input type="radio"/>
* To retrieve the document	<input type="radio"/>	<input checked="" type="radio"/>	<input type="radio"/>

\* 4.4 What is your perception of the response time?

- ☐ Good  
☒ Acceptable  
☐ Not acceptable

## 5 Patient Information

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### 5.1 Patient name and other information

5.1.1 Family name of the patient

PETERS

5.1.2 Prefix of the patient, if present

NA

5.1.3 Given name of the patient

Norbert Claude

5.1.4 Gender of the patient, if present

- ☒ Male  
☐ Female  
☐ Undifferentiated  
☐ Null value

5.1.5 Date of birth of the patient

01/01/1937

### 5.2 Patient identifiers

5.2.1 Regional/National Health Id

2544557646 (1.3.182.2.4.2), 1937010112347 (1.3.182.4.4)

#### 5.2.2 Social/Insurance Number

#### 5.2.3 Please, introduce here any comment you consider relevant regarding patient identifiers

Unable to identify the social/ insurance number

## 6 Original Clinical Document List

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\* 6.1 Are you able to select the specific Original Clinical Document to be accessed by looking only to the Original Clinical Document list information?

- ☒ Yes  
☐ No

\* 6.3 Did you encounter problems opening this or any other Original Clinical Document?

- ☐ Yes  
☒ No

## 7 Original Clinical Document

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7.1 Considering the information provided as a whole, do you find the service useful for the care you intend to provide?

- ☐ Yes  
☒ No

7.2 Please introduce here any comment you consider relevant regarding the question above

The information needs to be translated. If it were, then our answer would have been different.

## 8 Additional Feedback

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Please, upload here additional feedback about the evaluation: ideally, the CDA document (XML file) that was displayed to you and the rendering of the CDA Display Tool (PDF file). That information will greatly help Solution Provider when evaluating the submission.

The maximum upload limit is 1 MB.

8.1 Please upload your file here:

The maximum file size is 1 MB

**bfde8e65-afdd-43c5-b754-fa65b1b92b80/2025-03-24T12-14-03.344427Z\_CDA\_EHDSI---PIVOT-CDA-\_L1\_-  
VALIDATION---WAVE-8-\_V8.1.0\_\_NOT-TESTED.xml**  
**141e7365-e5b0-4a65-8cd4-f5a281f46df4/2025-03-24T13-02-49.249266Z\_CDA\_EHDSI---PIVOT-CDA-\_L1\_-  
VALIDATION---WAVE-8-\_V8.1.0\_\_NOT-TESTED.xml**  
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3.182.2.4.6.6.1.4274553371830469638\_lr.pdf**  
**ff711cdd-cb26-435c-a6ae-87f0dab31742/Binary\_urn\_oid\_1.3.182.2.12.10000734.6.1\_1.3.182.2.4.6.2\_1.  
3.182.2.4.6.6.1.5782983065764577735\_hd\_\_1\_.pdf**  
**58c2d9d6-6b06-42c5-9bdc-8803def1d846/LU\_L1\_OrCD\_Hospital\_Discharge\_04\_25.htm**  
**72bc620c-0d1d-4740-8e14-88c413082842/LU\_L1\_OrCD\_Laboratory\_Result\_04\_25.htm**

## Contact

[Contact Form](#)