

# CEN IPS Final meeting

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***Conformity Assessment of IPS implementation***

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# Conformity assessment for IPS

- IPS standard provides an abstract definition of a Patient Summary from which derived models are implementable
- A patient summary implementation shall be an IPS Document, comprising six mandatory IPS Data Blocks

- Patient Attributes
- Allergies and Intolerances
- Medication Summary
- Problems
- Provenance
- Cross Border (conditional)
- With same purpose
- Fulfil the conformance rules

Functional status  
History of past illness  
History of pregnancy  
History of procedures  
Immunization  
Medical devices  
Plan of care  
Results  
Social history

# IPS - Various implementations

- Country and/or cross border specific
- Various use cases (various actors including the patient): unscheduled/scheduled care, cross border/national/local, chronic patient,...)
- Various terminologies
- Standard implementation: HL7 CDA, FHIR, others
- ...



How to assess the conformity of  
the IPS implementation ?

# EURO-CAS key goal

*The goal of EURO-CAS is to develop the sustainable Conformity Assessment Scheme for Europe (CASforEU), to promote the adoption and take-up of interoperability testing of eHealth solutions against identified eHealth standards and profiles defined in the Refined eHealth European Interoperability Framework*

# EUROCCAS



# Pragmatic approach... and now to work!

- CASforEU defines a governance and organisation and processes for
  - Selecting of the test platform
  - Defining test cases
  - Selecting test methods (test tools and test data) and its extensions
  - Selecting and autorizing ISO/IEC 17027 test labs
  - Reporting tests performed by SUTs
- Existing tools that can be adapted for assessing various IPS implementation:
  - Art Decor
  - Gazelle Test management: support test cases and test workflow and management
  - EVS Client and Object checker
  - FHIR tools, Simplifier
  - Proxies, simulators, ...Ex: Test tools used in eHDSI
- CASforEU provides implementation guideline for various types of stakeholders (country, vendors, test laboratories, implementation entities, national and regional authorities or projects, etc

# EURO-CAS



# Benefits

## Vendors and IT companies

- Reduced effort in interoperability testing
- Investments redirected to innovative product features and usability
- Broadened market opportunities in a European Digital Single Market (and beyond)
- New opportunities for test laboratories

## Professionals, patients

- Quality, interoperability, usability of solutions
- Better time-to-market of innovative solutions
- Enhanced patient's engagement and mobility through such innovative solutions
- More resources, directly or indirectly, redirected towards the patient

## eHealth Initiatives, governments, policy makers, procurers, payers

- An independent benchmark
- Reduced effort and expenses in specification and testing
- Conformity with regulation

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# Questions?

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