

Building Personalized medicine In Switzer land



Christian Lovis MD MPH

çampus

biotech





NATIONAL REGULATORY FRAMEWORK – CARE SYSTEM FEDERAL LAW FOR THE SHARED PATIENT RECORD - 2017

Ablauf der Referendumsfrist: 8. Oktober 2015

Bundesgesetz über das elektronische Patientendossier (EPDG)

vom 19. Juni 2015

Die Bundesversammlung der Schweizerischen Eidgenossenschaft, gestützt auf die Artikel 95 Absatz 1 und 122 Absatz 1 der Bundesverfassung¹, nach Einsicht in die Botschaft des Bundesrats vom 29. Mai 2013², beschliesst:

1. Abschnitt: Allgemeine Bestimmungen

Art. 1 Gegenstand und Zweck

¹ Dieses Gesetz regelt die Voraussetzungen für die Bearbeitung der Daten des elektronischen Patientendossiers.

² Es legt die Massnahmen fest, die die Einführung, Verbreitung und Weiterentwicklung des elektronischen Patientendossiers unterstützen.

³ Mit dem elektronischen Patientendossier sollen die Qualität der medizinischen Behandlung gestärkt, die Behandlungsprozesse verbessert, die Patientensicherheit erhöht und die Effizienz des Gesundheitssystems gesteigert sowie die Gesundheitskompetenz der Patientinnen und Patienten gefördert werden.

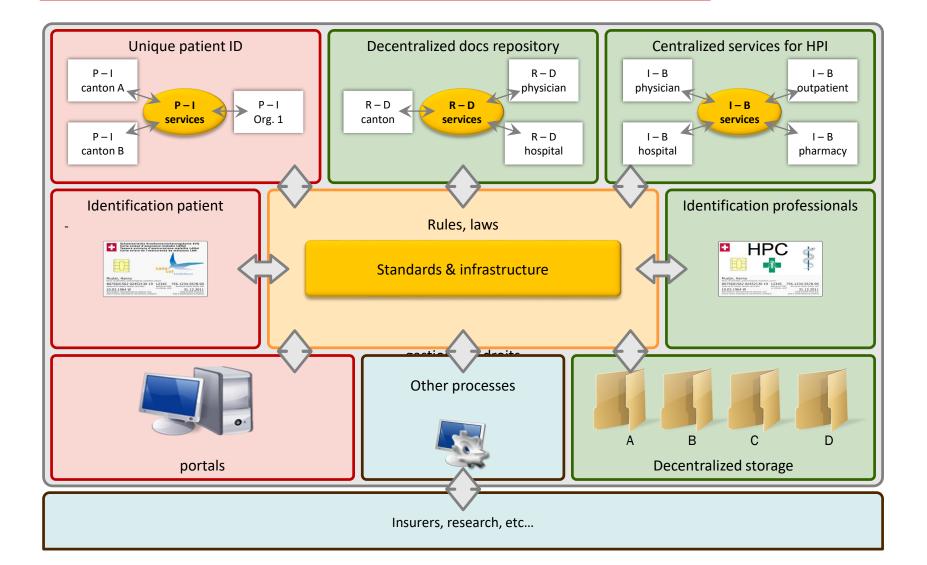
⁴ Die Haftung der Gemeinschaften, der Stammgemeinschaften, der Portale für den Zugang der Patientinnen und Patienten zu ihren Daten (Zugangsportale), der Herausgeber von Identifikationsmitteln, der Gesundheitsfachpersonen sowie der Patientinnen und Patienten richtet sich nach den auf sie anwendbaren Vorschriften.

EPDG - 2017 V1

DATENSCHUTZ GESETZ V2

DRAFT EID LAW V1

NATIONAL REGULATORY FRAMEWORK – CARE SYSTEM DISTRIBUTED GRID OF CERTIFIED COMMUNITIES OF CARE PROVIDERS



NATIONAL REGULATORY FRAMEWORK – CARE SYSTEM INTERNATIONAL STANDARDS OF INTEROPERABILITY

IHE driven process and methodology

- IHE XCA-XDS profiles
- HL7- CDA data models
- LOINC and SNOMED CT semantics

Swiss Personalized Health Network Initiative 2017 - 2020

2014	Concept
------	---------

2015 Plan

2017 Start

2016 Structure

- "Personalized Health". Report to discuss options and frame for a new Swiss Initiative.
 By an interdisciplinary and inter-institutional working group, June 2014.
- 2) Konzept Nationale Förderinitiative: "Systems Medicine Personalized Health".
 1st SERI commissioned Report by Working Group 1, September 2014.
- Interoperability of clinical and omics data in Switzerland
 2nd SERI commissioned Report by Working Group 2, February 2015
- 4) Setting up an Organisational Structure until end of 2015
 - 3rd SERI commissioned Working Group 3

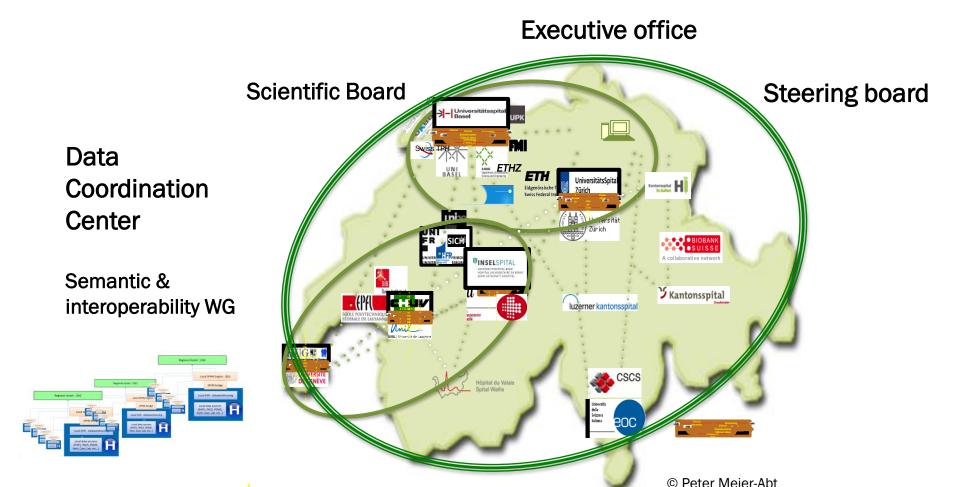


Akademien der Wissenschaften Schweiz Académies suisses des sciences Accademie svizzere delle scienze Academias svizras da las scienzas Swiss Academies of Arts and Sciences

© Peter Meier-Abt

SWISS PERSONALIZED HEALTH NETWORK INITIATIVE DISTRIBUTED NETWORK OF REGIONAL CLINICAL DATA CLUSTERS

Swiss Personal Health Private Research Cloud - SPH-PRC



I2B2/TRANSMART FOUNDATIONS STANDARDS, SHRINE DISTRIBUTED NETWORK OF REGIONAL CLINICAL DATA CLUSTERS

Swiss Personal Health Private Research Cloud - SPH-PRC

i2b2 Informatics for Integrating Biology & the Bedside

About Us | Software | NLP Data Sets | i2b2 tranSMART Foundation |

Collaborations

- Scientific Collaborations
- i2b2 Installations
- Academic Users' Group
- Data Sharing Network (SHRINE)
- Use of our Software
- Data Sharing Policy

Data Sharing Network (SHRINE)

In their application for CTSA status, Harvard Medical School (HMS) proposed creating a webbased sotware network (based on the prototype model SPIN, Shared Pathology Informatics Network) that would allow the participating Harvard hospitals to link their respective i2b2 instances for the sharing of obfuscated, aggregated counts of patients meeting selected inclusion and exclusion criteria for demographics, diagnoses, medications, and labs. It was envisioned that this network, called SHRINE (Shared Health Research Informatics NEtwork), would greatly enable population-based research, assessment of potential clinical trials cohorts, and hypothesis formation for followup study by combining the EHR assets across the hospital system.

A National Center for Biomedical Computing

SPHN SMARTCONSENT



CITIZEN CENTERED CONSENT (C3) MANAGEMENT SYSTEM



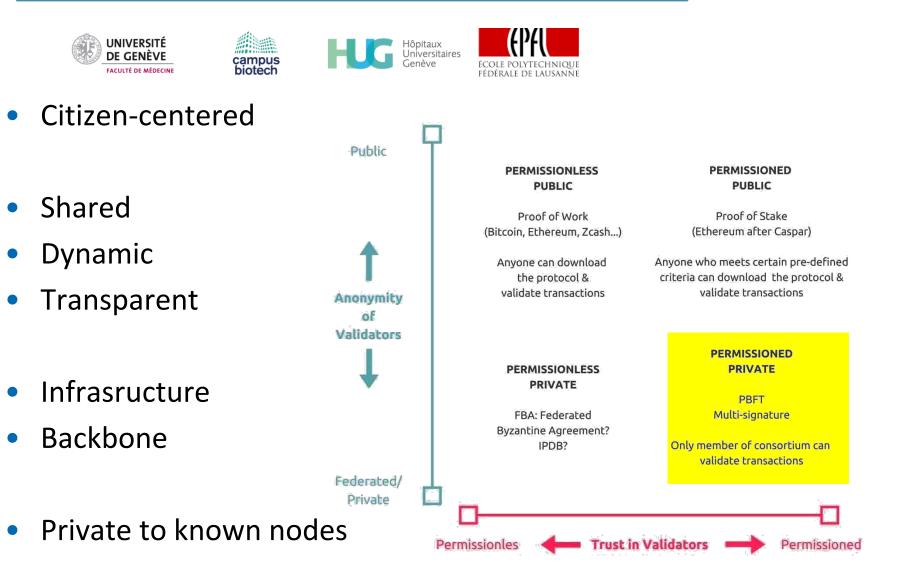




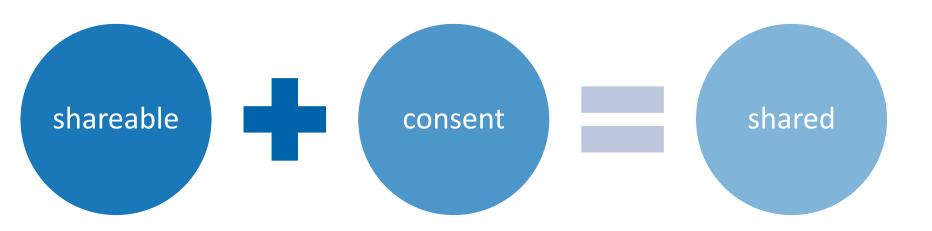


- Blockchain based smart contracts
- Compositional model of the consent
- Specific implementation of a contract with an indefinite parts
- RESTful API implementation of the service
- In test currently
- 6 trusted distributed authorities foreseen:
 - 5 CH university hospitals
 - SwissEthics

SPHN SMARTCONSENT CITIZEN CENTERED CONSENT (C3) MANAGEMENT SYSTEM



SWISS PERSONALIZED HEALTH NETWORK INITIATIVE DATA STRATEGY



Interoperability Clear model Strong semantic Rich metadata Generic model Citizen centered «all consents» One cohort in CH:

All citizen

Smartcontracts Dynamic shared Protection by design «fit-for-purpose» sub-cohort management «quality management