

What the Plan and Framework mean for Patient Care

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Discussion points



- Brief Current State
- Policy data needs
- Information Access
- Care Access
- Quality and Patient Safety
- Building Trust and Confidence



Current Data Ecosystem

It can still be difficult to share health data, even in emergencies. This is because health stakeholders such as hospitals, general practitioners or researchers collect data in many different formats which are incompatible with one another.

- Paper charts (multiple)
- Electronic systems
- Uncertain identity
- Data Silos
- Linkage capability very low



Reduce complexity and simplify

10 systems to get to here



Healthcare professionals along the patient journey



Integrated information for integrated care



Value Based Healthcare



Traditional Care

- Organized around specialties and interventions
- Duplication across sites/facilities
- Sites provide care for multiple acuity levels
- Limited Integration
- Siloed Data and IT systems

The current health service environment

<u>Slaintecare</u>

- Multidisciplinary teams managing full care cycle with the <u>patient</u>
- Clinically Integrated Care Delivery System
- Allocate services to sites based on complexity, risk and patient convenience
- Incorporate all modalities of care
- Use Formal systems to allocate patient care to most appropriate site

Dependency – Enabling Health Data & Information



Policy - Slaintecare

- "A healthier Ireland, with improved health and wellbeing for all, and with the right care delivered in the right place at the right time."
- Key enablers -data, technology, and digitisation,
 - "advancement in health systems information, is vital in underpinning policymaking that is person-centred and delivers on strategic priorities".

DOH Statement of Strategy 2021-2023

- Integrated care is only possible with free flow of Data and Information
 - Right Data and the Right Staff.



Inflection Point - Data & Information



"There comes a point where we need to stop just pulling people out of the river. We need to go upstream and find out why they're falling in"

-- Archbishop Desmand Tutu

Single Vision



Better Health Outcomes Enabled By

Seamless, Safe, Secure & Connected Digital Health Services

Supporting the Health & Wellbeing for

Both Patients & Providers



The HSE and the Department of Health have collaborated together to bring this vision to life.

Where will we Impact patient Care?



Enhanced Access to Information

Managed Health Information

- Comprehensive service Information
- Accurate sign posting
- Clinically approved health guidance
- Self-assessment and self-management tools
- Automatic and people based communication channels
- Targeted messaging –health promotion

Personal Health Data & Information

- Access to our own healthcare record
 - The data about me used and shared between the healthcare professionals who take care of me,
- Input to our own health data
- Raise concerns if we see errors
- Direct access to our appointment schedule
- Make/cancel appointments
- Interact directly with the care team



Improved Access to Care

Access to care options:

- Available health record
- Use staff to full scope
- More appointment slots
- Closer to home
- Virtual Care in your own home

Access to consultation options:

- In-person
- Virtual consultations
- Combined in-person and virtual with specialist
- MDT consultations directing appropriate care and avoiding duplication

Access to marginalized communities

- Support care at centres people attend with mobile devices health checks, vaccinations, etc.
- Language and translation supports through digital tools

Improved Access to remote communities

- Telehealth consultations/combined consultations
- Remote monitoring centres
- Online therapies

Barriers:

Digital Literacy

Health literacy

Privacy Concerns

Social circumstances

Principles 1, 2, 3, 4

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Patient Centred Care

Access to the single patient Record

- Stops repetitive requests for the patients story
- Patients who are ill often not able to provide the full story
- Full access to the medical chart:
 - Ensures earlier diagnosis
 - Leads to earlier treatment
 - Reduction in duplication
 - Better value care



Collaborative care

Paternalistic care Traditional care



Patient – centred care Integrated care

Achieved through:

- Empowerment
- Self-management
- Personalization of care



Personalized Care

- Preventative Health and Wellbeing
- Personalized care and therapies
- Disease prediction and prevention
- Genetics and genomics
 - Personalized treatments
 - Impacting rare diseases diagnosis, treatment, quality of life
- Precision medicine



Academic Health Service – EHR and Remote monitoring

•	Reduction in ED visits	25%	
•	Reductions in admissions	38%	
•	Decrease in costs	17%	
•	Preventable re-admissions	52%	
•	Medication compliance Earlier diagnosis/detection	13%	
	of recurrence	22%	

- Empowerment and Meds compliance
- Empowerment and disease management

89% relationship

80% relationship



Avoidable patient Harm (1:20)

- Medication Errors (50%)
- Patient misidentification
- Unsafe surgical procedures
- Diagnostic errors
- Patient falls
- Unsafe Blood transfusion
- Venous thromboembolism
- Pressure ulcers



Patient Safety – World Health Organization (WHO)

HE

Quality and Patient Safety

- Accurate identity
 - Right information right patient
 - Reduced data breeches
- Improved communication
- Significant clinical risk early identification:
 - Sepsis
 - Imminent collapse
 - Risk of falls
- Reduce medication errors
 - Allergy alerts
 - Drug interactions



Trust and Confidence

Through:

- A trusted, secure health and care ecosystem where data in managed, shared and stored securely, consistently and transparently
- Full rights to their data for patients/citizens involved partner in their health
- Consent around data access and use for data owners
- Protection of sensitive health data
- Protection of data rights with evidence of compliance
- Transparency on data use though audit, including anonymized data



Participatory Design

A collaborative design approach that involves end-users in the design process. Its aim is to create products and services that better meet the needs and expectations of users by applying their knowledge and experiences.



The key areas of consensus - IPPOSI citizens jury (2021):

We need a connected, quality, digital health information system

We need citizens to be the owners of their own information

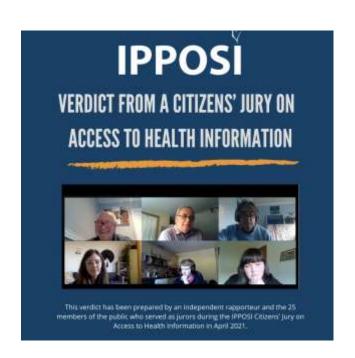
We have a collective responsibility to do the right thing

We need to be supported to grow trust and confidence in the State

We need to partner with citizens to design our health information future

We need to treat data as a national resource

We have to make consent the cornerstone of everything we do





Health Service Executive

Digital Health Strategic Implementation Plan 2024-2030

Thank you for listening

