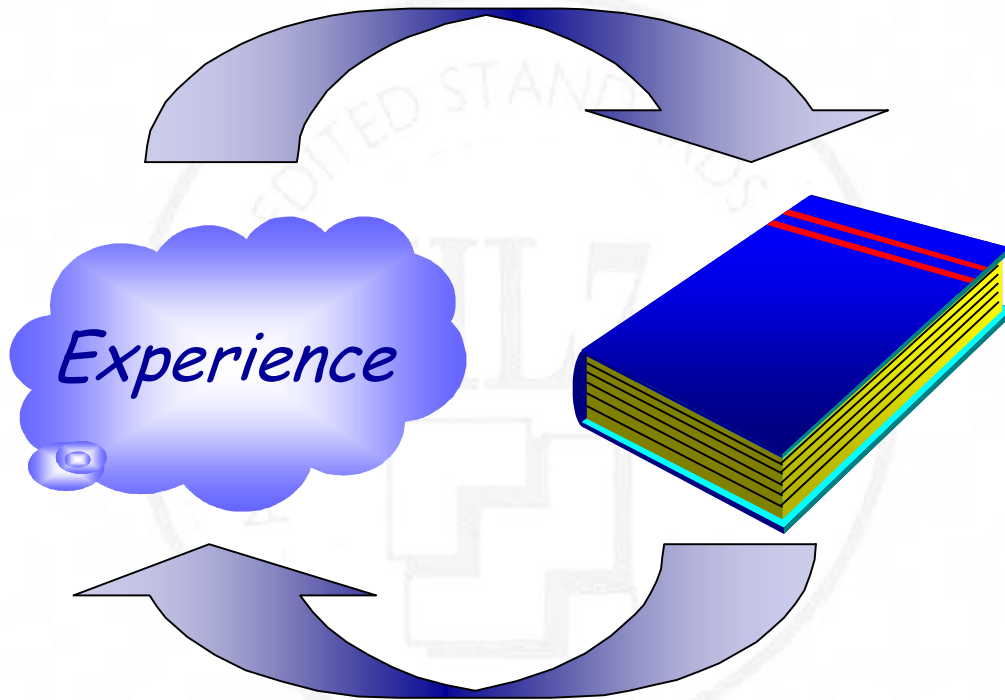


# **Message Formats, Models, and Syntax**

**Testimony for the National Committee  
on Vital & Health Statistics  
29 March 1999**

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# Standards/Experience Cycle



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1

## Versions 2.x

### Strengths

- ✓ broad functional coverage
- ✓ highly adaptable
  - IS environments differ
  - system capabilities variations
- ✓ vocabulary independent
- ✓ *least common denominator* technological base

### Difficulties

- ✓ broad functional coverage
- ✓ highly adaptable
  - “Seen one? Seen one.”
  - vendor capability mismatch
- ✓ vocabulary independent
- ✓ *least common denominator* technological base

## Functions of Comparable PMRI

- Extending the efficiency and efficacy of individual providers and organizations
- Extending the efficiency and efficacy of healthcare

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Extending the efficiency and efficacy of individual providers and organizations

availability of current information at the point of care

across provider organizations

within rapidly changing provider organizations

decision support

support for standards of assessment and care

support for the somewhat bewildering array of administrative constraints on decision making and requirements for supporting data

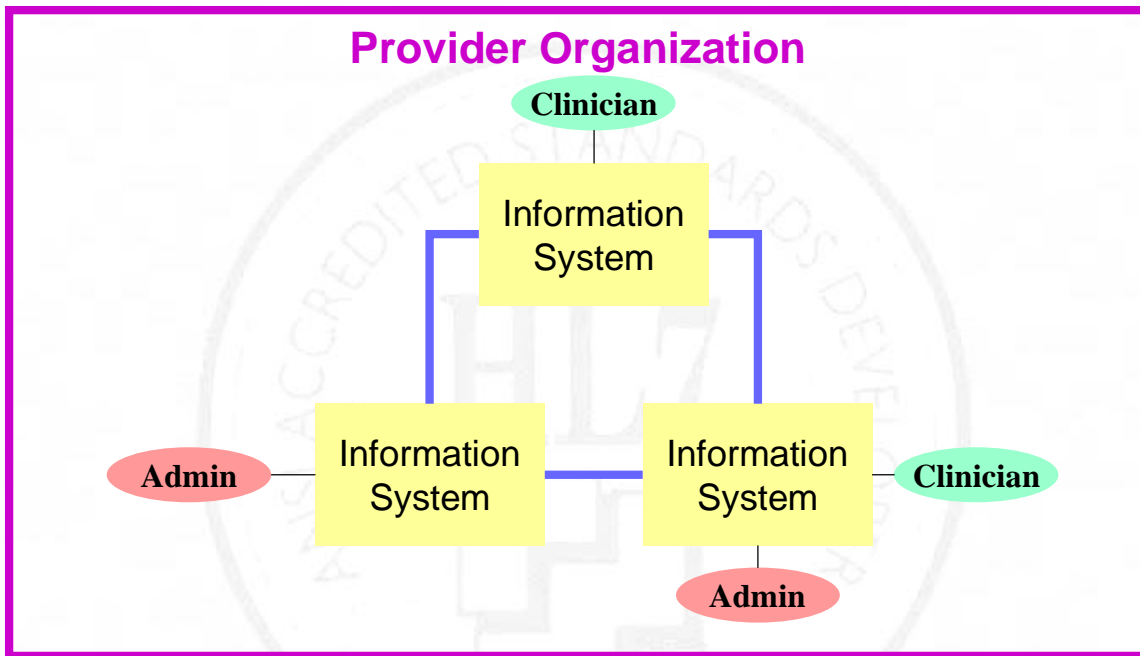
Extending the efficiency and efficacy of healthcare

evidence-based medicine

public health

public policy

# Interfaces Within an Organization

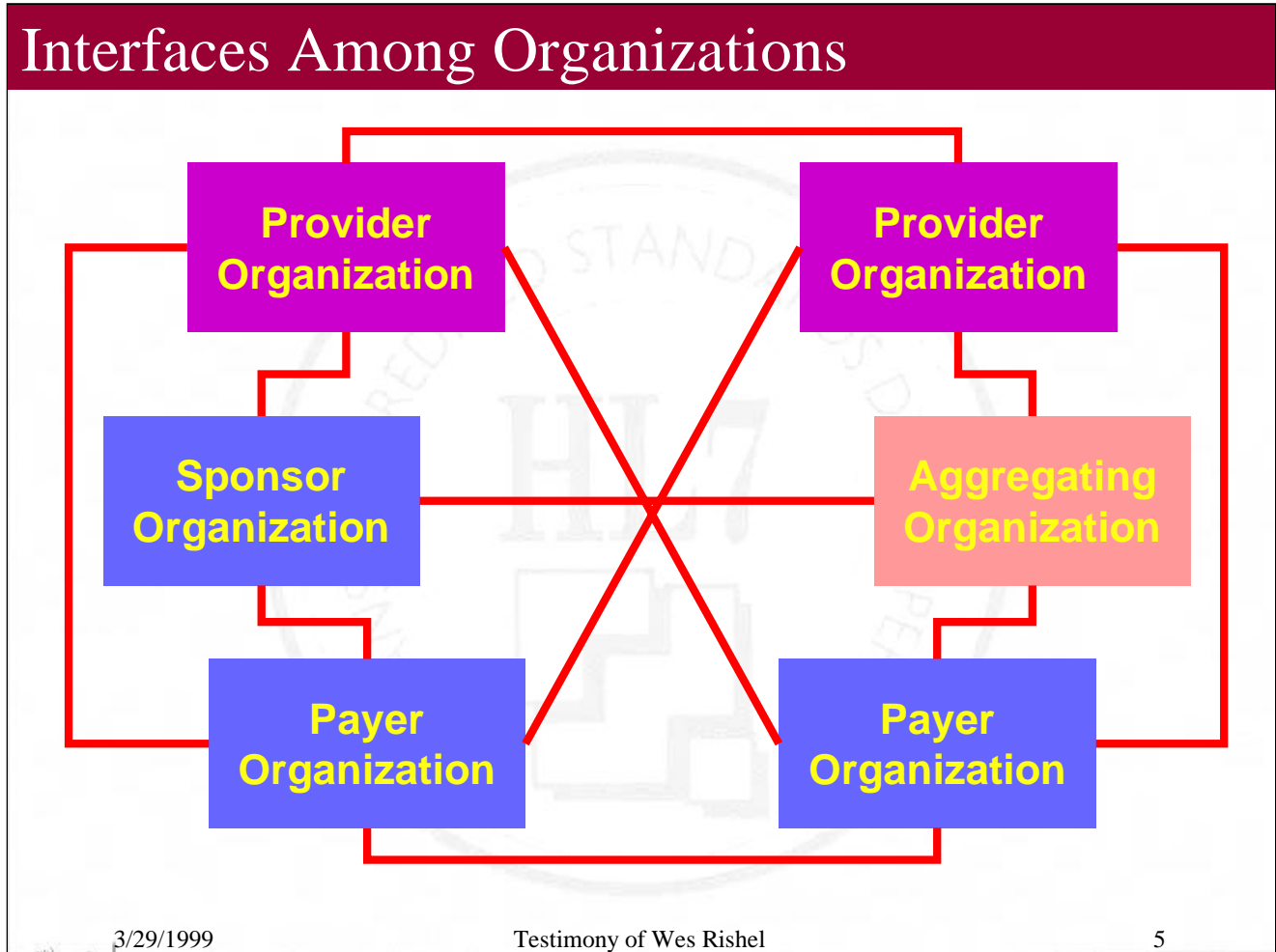


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Governmental or societal interest less direct  
assurance of quality  
assurance of availability of information



Governmental or societal interest is direct  
cost containment with quality  
assurance of availability of information  
barriers more difficult without government intervention

## Prerequisites for Exchanging Comparable PMRI



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Technologies and syntaxes are important, but the primary challenges are in specifying information content.

Technology and syntax are interrelated

Appropriate combinations of technology and syntax reduce the cost and other barriers to acceptance within the industry

e.g.,

reduced programming, debugging, and customization costs  
use of the Internet with suitable safeguards for  
interorganizational information transfer

## How Comparable is Comparable?

- What does it mean *not* to be comparable?
  - ✓ Communicate less information?
  - ✓ Communicate less precisely?
  - ✓ Tolerate a percentage of error?
    - acceptable for statistical applications
  - ✓ Have bilateral implementation agreements
    - costs measured on a “per system” basis
    - inconsistent with previous HIPAA requirements

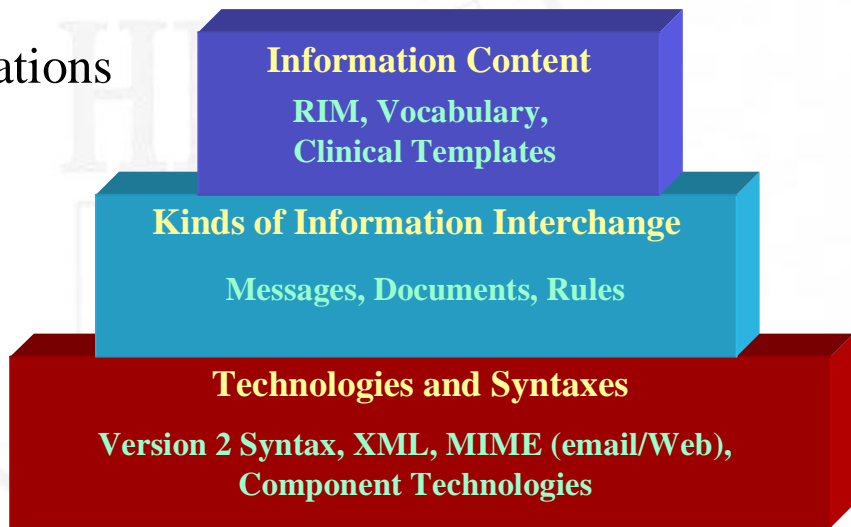


## What Does Comparability Cost?

- There is now, and for the foreseeable future, a cost barrier to full exchange of comparable fine-grained information
- The barriers are not primarily attributable to the syntax or technologies of information exchange
- They are attributable to:
  - ✓ the need for combined standardization on
    - information structure
    - vocabulary
- The cost of initially collecting the information to be exchanged

# HL7 Contributions

- Information Structures
  - ✓ Reference Information Model
  - ✓ Clinical Templates
  - ✓ Semi-definitive approach to Vocabulary
- Expanded Applications
- All Appropriate Technologies



## The Approach

- **The HL7 Vocabulary TC is committed to using existing vocabularies (coding systems) as values for coded fields in HL7 messages, rather than creating a new terminology.**
- **We need a solution that allows HL7 to reference and use proprietary vocabularies (SNOMED, Read, etc.) in a manner that is equitable to all vocabulary creation/maintenance organizations.**

## HL7 Assumptions

- Should **not** choose a single proprietary scheme
- Free-for-use coding systems preferred
- “Market model” will assure responsive maintenance
- Proprietary vocabulary use requires a license

# ORU: The usual result message

<b>ORU</b>	<b>Observational Results (Unsolicited)</b>
<b>MSH</b>	<b>Message Header</b>
<b>PID</b>	<b>Patient Identification</b>
<b>{OBR</b>	<b>Observations Report ID</b>
<b>{OBX}</b>	<b>Observation/Result</b>
<b>}</b>	<b>Answer Part Loop</b>
	<b>Element Loop</b>

## How much meaning to put in a code?

### HL7 OBX Segment (simplified)

**OBX | Observation Id | Value | Units**

**OBX | 123^Ser Na Conc | 138 | mmol/L**

### Possible Codes for Observation Id

**123 Serum Na Concentration**

**456 Serum Na Concentration (mmol/L)**

Code meanings must be consistent with the data structure within which they will be used !

## The Missing Link: Clinical Templates

- Universal “trading partner agreements”
- More specific
- More costly

## An instance of an ORU message

**MSH, EVN, PID, PV1, ORC,**  
**OBR||8974-9^BP Battery^LN|**  
**OBX|1|CE|8357-6^METHOD^LN|M^Manual|**  
**OBX|2|CE|8358-4^DEVICE^LN|1|AC^Adult Cuff|**  
**OBX|3|CE|8359-2^SITE^LN|1|RBA^Rt Brachial Artery|**  
**OBX|4|CE|8361-7^POSITION^LN|1|SIT^Sitting|**  
**OBX|5|NM|8479-8^SBP^LN|1|138|mmHg|**  
**OBX|6|NM|8462-4^DBP^LN|1|85|mmHg|**



## A BP Battery Template

### Battery Level Constraint

```
BPBattery ::= SET {  
  obr {  
    universalServiceID (8974-9^BP Battery) }  
  obxs {  
    MethodObs,  
    DeviceObs,  
    SiteObs,  
    PositionObs,  
    SystolicBPObs,  
    DiastolicBPObs }  
}
```

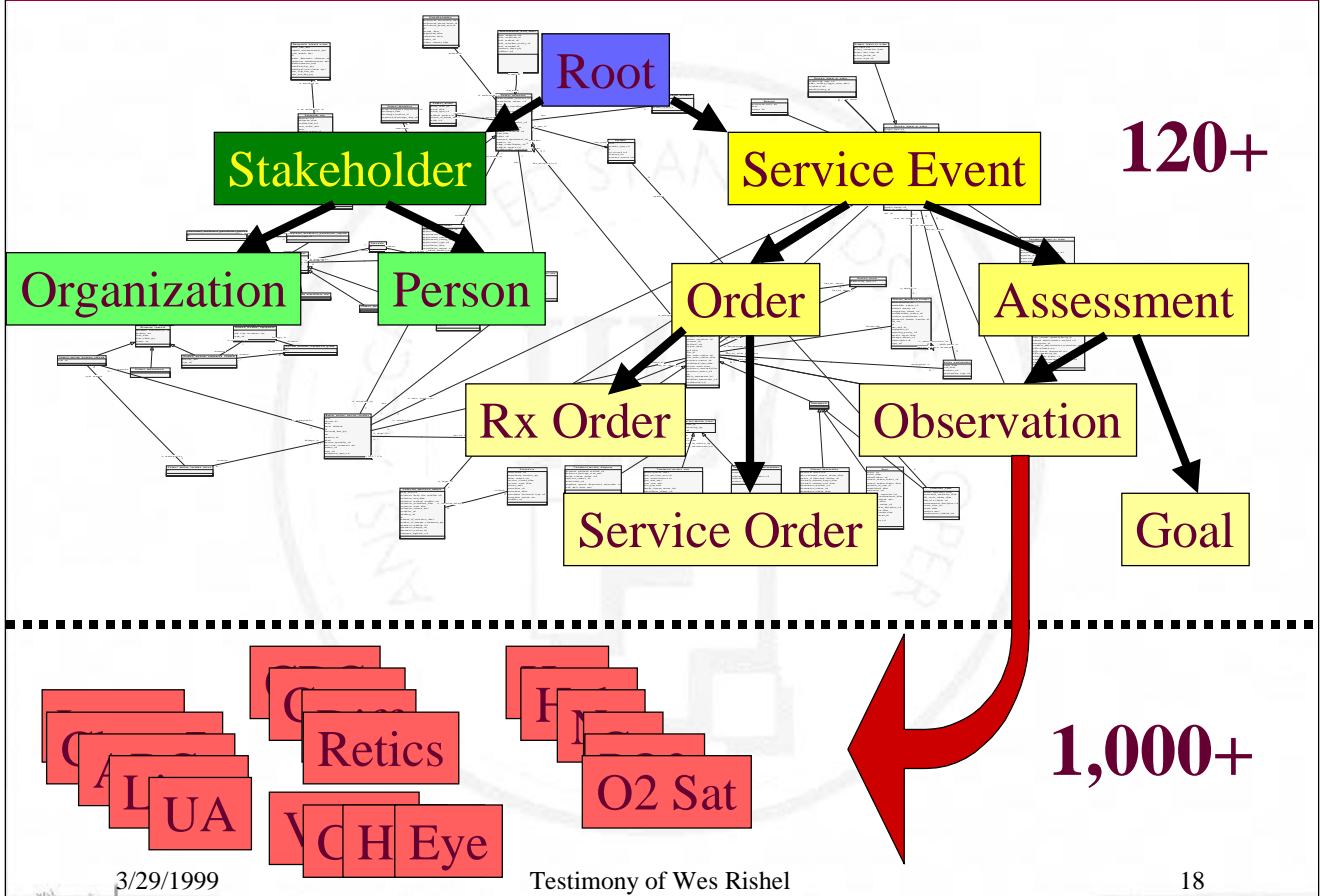
## Observation Template

### Observation level constraint

**PositionObs ::= SET{**  
    observationId (8361^POSITION^LN),  
    value (PositionDomain) }

**SystolicBPObs ::= SET{**  
    observationId (8479-8^SBP^LN),  
    value (Numeric, “DDD” ),  
    units (mmHg) }

# The RIM and Clinical Templates



## What are clinical templates?

- “*Constraint of an existing information model*”
  - ✓ Reference Information Model (RIM)
- Constraint of specific RIM classes:
  - ✓ clinical assessment
  - ✓ clinical observation
  - ✓ service event
  - ✓ service order
  - ✓ others
- Can be applied to Version 2.X or version 3.0

## What Kinds of Data Will Templates Describe?

- **Clinical laboratory batteries**
- **Physical exam findings**
- **Microbiology culture results**
- **Immunization queries and results**
- **Claims attachments**
- **Medical Records Documents (transcription)**
- **And hundreds more ....**

## Clinical Template Work Items

- **Define the formal notation for templates**
- **Define a process for creating, approving, and maintaining templates**
- **Define a mechanism for participating with professional societies or other clinical experts in the creation of template content**
- **Create a repository for storing and allowing access to templates**

## Where Will Clinical Templates be Used?

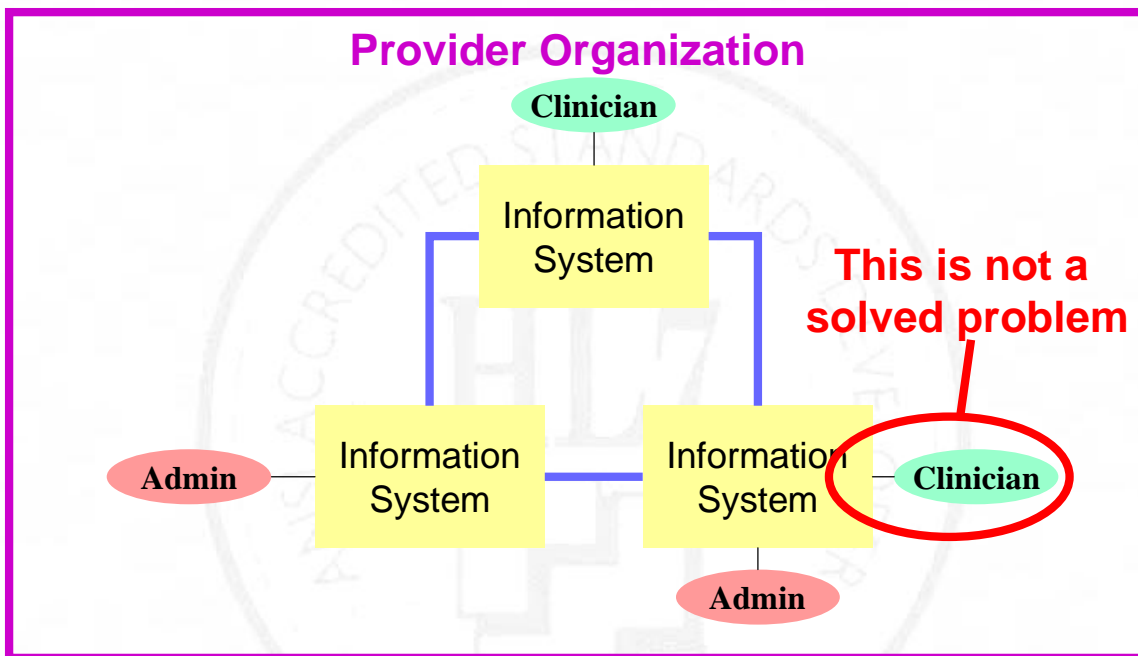
- **Messages with fine-grained, comparable, clinical data**
- **Documents**
- **Rules**

## HL7-HIPAA Claims Attachments

- Collaborative team: HCFA, X12, HL7
- HL7 version 2.3 syntax embedded within X12 275 transaction
  - ✓ Rigid pattern of segments
  - ✓ Variability through LOINC codes
- No bilateral trading partner agreements
- Varying degrees of granularity
- Comparability requires more or less human judgement
- This is a primitive form of clinical template



# The Other Cost of Fine-Grained Information



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Much of the most valuable information comes from providers making care decisions

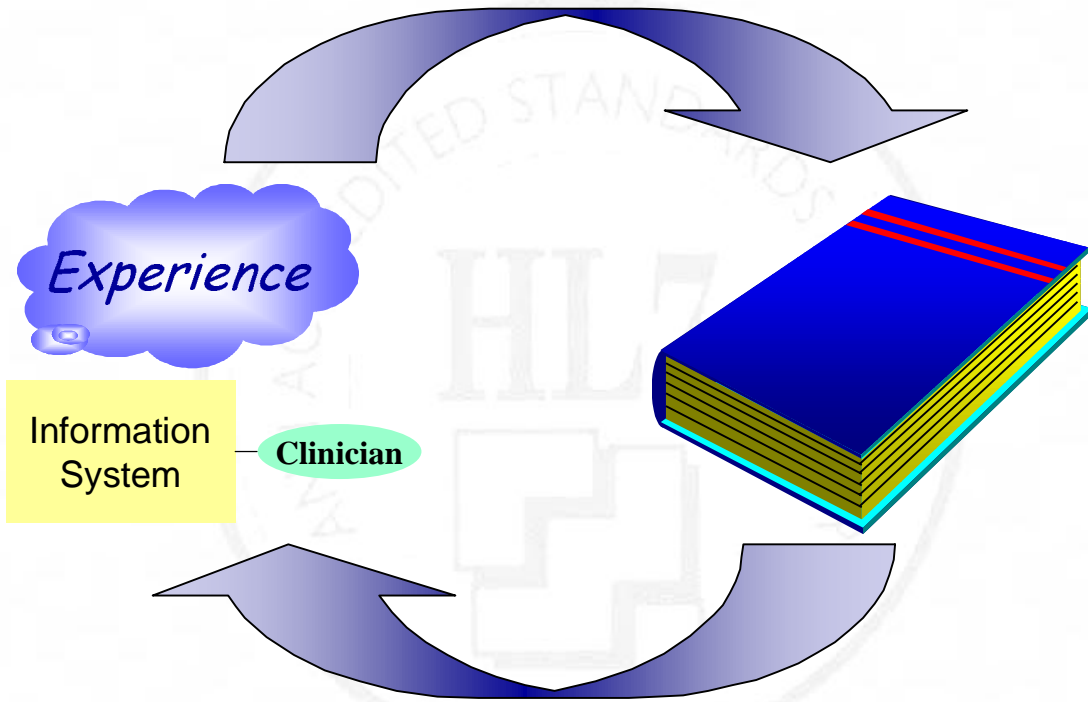
Capturing information requires provider interaction with the computer at or near the time of making the decisions

The time of the provider is itself a major cost

We rely on medical record designs to co-optimize physician time and information quality

This is not a solved problem

## Co-Evolution of EMRs and Standards



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The issues are not wholly separable

The latency time for improvements is substantial

- standards development

- system engineering cycles

- provider implementation cycles

The situation has similarities to automobile mileage and pollution regulation

## What Can the Government Do?

- Establish Time-Based Goals With Teeth
- Require no trading partner agreements for aggregation requirements
- Don't let the perfect be the enemy of the good
  - ✓ **early on:** emphasize selective achievements consistent with current systems and extant standards
- Don't let the good be the enemy of the better
  - ✓ **long term:** *raise the bar* to promote finer-grained standards in time frames consistent with industry product cycles

## Specific Governmental Actions

- Support the creation of clinical templates
  - ✓ Methodology
  - ✓ Outreach to clinical specialty groups
- Strongly influence in the business of providing vocabularies
  - ✓ Select, sponsor, negotiate public rates for *a few* important vocabularies
    - Drugs
    - Signs, symptoms, diagnoses, history, physical exam
    - Laboratory findings
    - Observation identification