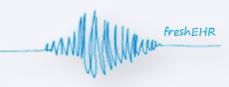
A new approach to clinical standards

Dr Ian McNicoll SCIMP Co-chair openEHR HANDIHealth CIC



SCIMP/SNUG conference Sept 2015

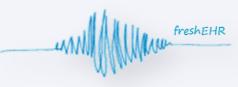






Current scenario

- Patient often only the knows the whole picture
- No clear governance
- Non- standardised representation of medication between systems
- No clear visibility of other prescribers actions



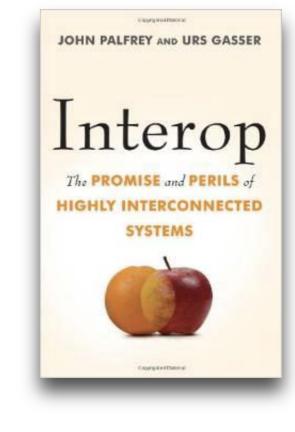




Interoperability is not a tech problem

"The real barriers to practical interoperability are cultural and clinical"

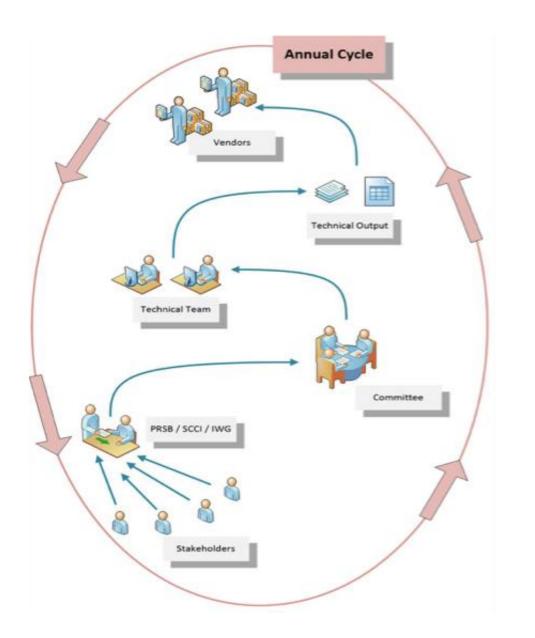
- Diverse recording practice (sometimes arbitrary)
- Diverse recording requirements
- Complexity / contextual nature of health data
- Lack of clinical involvement in standards development
 - Too technical, too philosophical
 - Too time-consuming, too slow





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Traditional standards development



- Clinical stakeholders engage through top-down governance
- Committee-based
- Late vendor engagement
- Fixed review cycles
- Unclear / unresponsive change request mechanism







Are 'Standards' necessary?

FAREWELL TO "RUTHLESS STANDARDISATION"

© SEPTEMBER 2, 2014 ▲ WOODCOTE PLEAVE A COMMENT

"Ruthless Standardisation" was the failed mantra of the NHS National Programme for IT. The Programme is dead, but in some places this view still persist but it is time to consign it to history as something else that "seemed a good idea at the time" http://www.woodcoteconsulting.com/farwellto-ruthlessstandardisation/

Are standards necessary?

November 1, 2013 § 9 Comments

THE GUIDE TO HEALTH INFORMATICS ENRICO COIERA

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A COMMON STRATEGY FOR STRUCTURING COMPLEX HUMAN SYSTEMS IS TO demand that everything be standards-based. The standards movement has taken hold in education and healthcare, and technical standards are seen as a prerequisite for information technology.

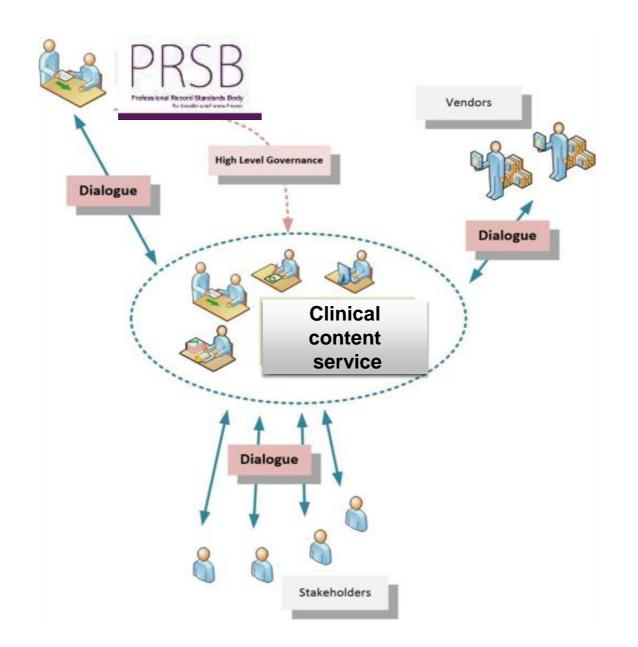
In healthcare, standards are visible in three critical areas, typical of many sectors:



http://coiera.com/201 3/11/01/arestandardsnecessary/

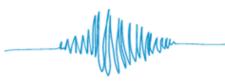


Clinically-led standards development



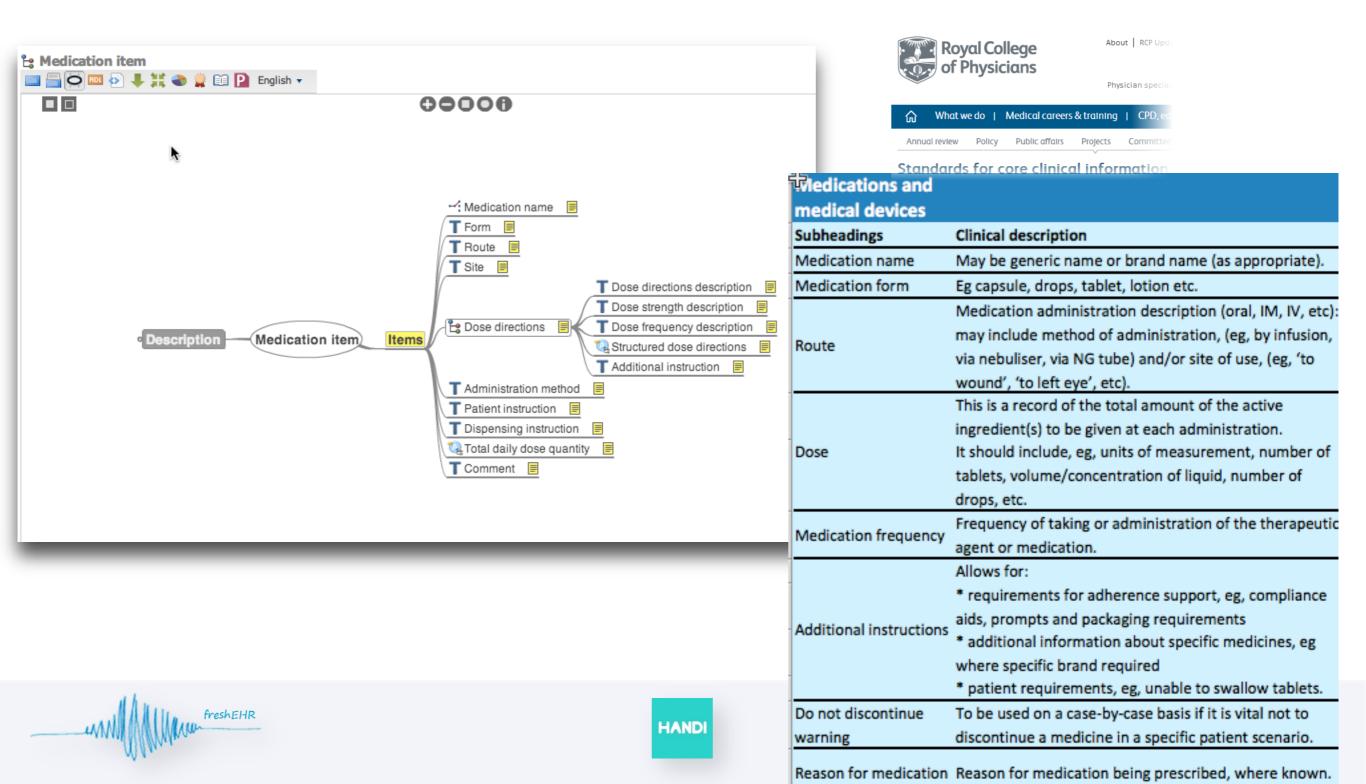
- Clinical stakeholders, vendors engage directly with clinically-led content service
- Continual dialogue with all stakeholders via web-based collaborative tooling
- No fixed review cycles
- On-demand change request directly to clinical content service
- PRSB has high-level governance role





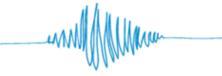


Medication 'archetype'



Web-based clinical collaboration

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Project or Incubator:					
•All active Under review Published	Purpose Not Specified				
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Incubators +					
Try New and modified Resources +	Date First Verified 11/05/2014				







Web-based clinical review

Content Review Summary: Adverse reaction

Switch to detailed view

Content Review Summary: Adverse reaction (Revision: 6) (! Sam Patel (03-May-2013)

Invitation

Headan

Co

Paul Miler (17-Apr-2013)

I stil feel the name is fundamentally misleading to clinicians as the archetype is to be used for adverse reactions, not just immune mediated reactions.

Heather Lesle (30-Apr-2013)

There is debate in many circles, but it can commonly be agreed that allergies and intolerances are a subset of the broder notion of an adverse reaction. On the other hand it is not clinically understood that intolerances are a subset of allergies, which is implied by the naming. So being devils advocate here - I recognise that naming this archetype is largely for historical reasons, but given that this archetype may be in use for many years to come, is it worth considering renaming the concept for posterity? Further, are you recording the allergy or evidence of the allergic reaction? There are many that argue that this is a really important distinction.

119511003]

between allergy and adverse reaction. The latter can fall within a side effect profile and be addressed if not severe.i.e. nausea. I understand that the latter fields may compensate for this , but unless it is clear, agents to treat Ife threatening conditions may be excluded because of an 'allergy' label. E.g patients with severe streptococcal infection will do better on a penicilin, but this may be excluded because of a vomiting episode on one occasion and the label of 'allergy/adverse reaction'. If there is sufficient detail in the remainder of the archetype then fine, but I feel you can't have one without the other.

I speak from a secondary care

boat here, but a distinction is vital

perspective and I will probably rock the

Colin Brown (29-Apr-2013)

prefer "Adverse Reaction" as a more inclusive term for titles etc, it includes "alergies". As implied by SCT's term I think BMJ articles supported this a few years ago - could search it out ...



Editor Feedback

@Paul @Coin I agree that adverse reaction would be a better term. This archetype has its roots in the GP2GP 'Drug alergy' archetype where its use is quite limited to the recording of drug allergies and adverse reactions

@Heather

g (dr

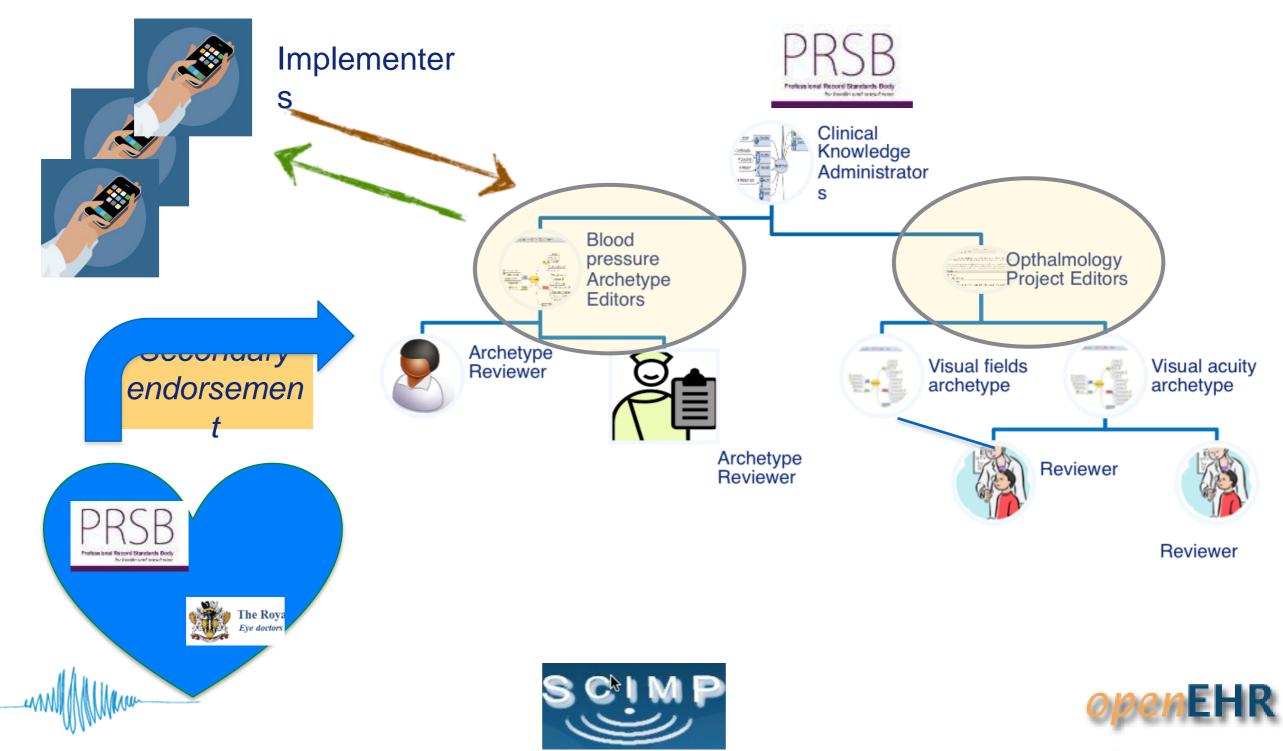
1) I agree that it is worth renaming to 'adverse reaction' in line with thinking elsewhere.

2) We are essentially recording the risk of / propensity to allergy/adverse reaction, with a single code for the reaction observed. UK GP systems all simply capture a single 'allergy' record which mixes the record of the reaction with the assertion of future risk. I know this is hotly debated around the world but at least in the GP systems community we have some real consensus in recording practice.

@Sam This is also one of the aspects that is a hot topic when the recording of allergy is discussed. Should we try to distinguish allergy from adverse reaction from intolerance etc? In practice it has been found that clinicians are pretty poor at making the distinction reliably and, particularly in general practice, the nature of the underlying pathophysiology can be pretty unclear.



Evolutionary standardisation 'distributed Governance'



openEHR Foundation openehr.org

Home	Programs	Getting Involved	Downloads	News & Events
What is openEHR? Who is using open openEHR Specification openEHR Clinical M	ations Models		Management Board Election Feb 2015 Nominate now! What is the Management Board? Board elections page Current nominations	Indust

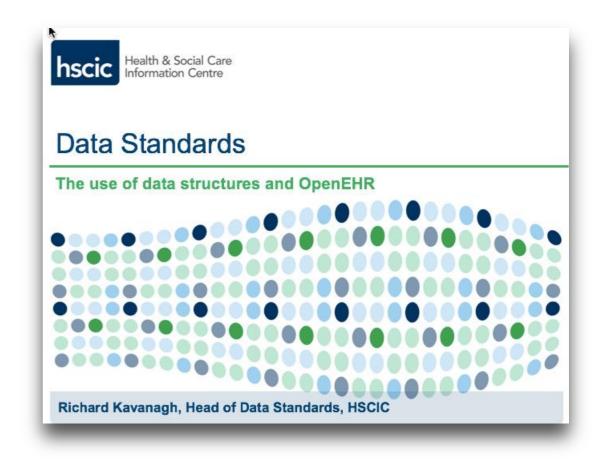






HSCIC

- have adopted openEHR as primary 'clinical content standards methodology'
- acquiring a separate 'English CKM'
- developing archetypes for
 - PRSB Transfer of Care summary
 - GPOC-R / GP2GP
 - Allergies, Blood pressure, Document metadata



http://www.infostandards.org/ documents/data-structuresv1-0-richard-kavanaghpptx.pptx



Professional Records Standards Body

- Professional Records Standard Body (4-country)
 - adopting NHS Scotland / GP2GP medication archetypes?
 - Relationship with HSCIC
 - HSCIC (largely) fund PRSB
 - PRSB commission work from HSCIC

00000		Wedications and medical devices	
		Subheadings	Clinical description
		Medication name	May be generic name or brand name (as appropriate).
		Medication form	Eg capsule, drops, tablet, lotion etc.
		Route	Medication administration description (oral, IM, IV, etc may include method of administration, (eg, by infusion via nebuliser, via NG tube) and/or site of use, (eg, 'to wound', 'to left eye', etc).
		Dose	This is a record of the total amount of the active ingredient(s) to be given at each administration. It should include, eg, units of measurement, number of the state of the state of th
	T Site		tablets, volume/concentration of liquid, number of drops, etc.
	T Dose directions description	Medication frequency	Frequency of taking or administration of the therapeut agent or medication.
	T Dose strength description		Allows for: * requirements for adherence support, eg, compliance aids, prompts and packaging requirements
(Medication item) Items	Additional instruction	Additional instructions	 additional information about specific medicines, eg where specific brand required
	Administration method		 patient requirements, eg, unable to swallow tablets.
		Do not discontinue	To be used on a case-by-case basis if it is vital not to
	T Patient instruction	warning	discontinue a medicine in a specific patient scenario.
	T Dispensing instruction	Reason for medication	Reason for medication being prescribed, where known
	Total daily dose quantity	1000	
	Comment		



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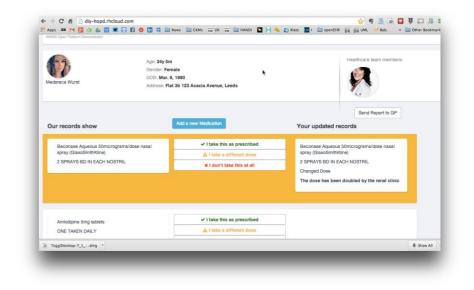
NHS England



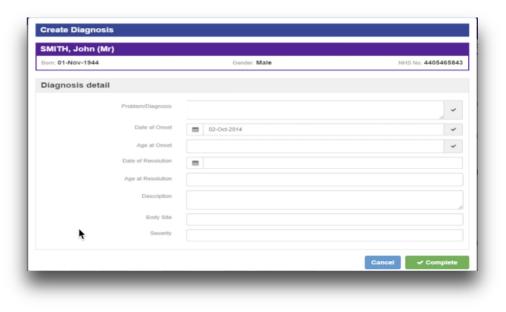
- Using EHRScape engine for SME / clinical training
- Supporting openEHR-based 'Open platform' approach
 - OPENeP

freshEHR

- Ripple OSI / openDental
- part share 'UK CKM' licence



Code4Health





NHS-E Interoperability Handbook

7.6 Data structures

Data structures are structured definitions of discrete data entries within care records that are commonl ensure that data entries that are determined to have the same logical meaning are represented consis flagged as either a **Standard**, as **Guidance** or as **Policy**.

Resource	Further information	Link to reso
OpenEHR	OpenEHR is a health record standard prevalent in research and used in provider systems in a number of countries. It utilises a reference model that extends the ISO 13606 reference model as well as archetypes to model care record data structures. These archetypes may then be constructed into templates (to represent datasets) which provide the basis for storing and exchanging electronic health records. A programme of work to collaborate with local organisations on defining data structures for national publication is currently underway within HSCIC. This includes work on the development of Discharge Summary interoperability standards.	http://www.openehr.org/



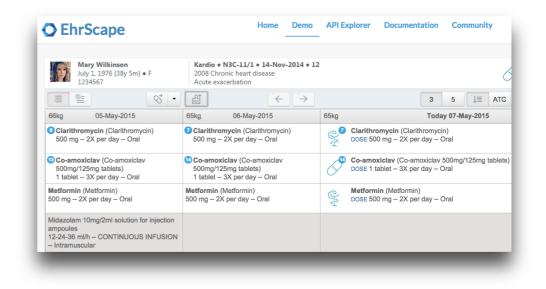
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OPENeP



- open-source HEPMA solution built on openEHR back-end
 - will use NHS Scotland medication archetypes for discharge and reconciliation
 - tendering for Scottish HEPMA framework

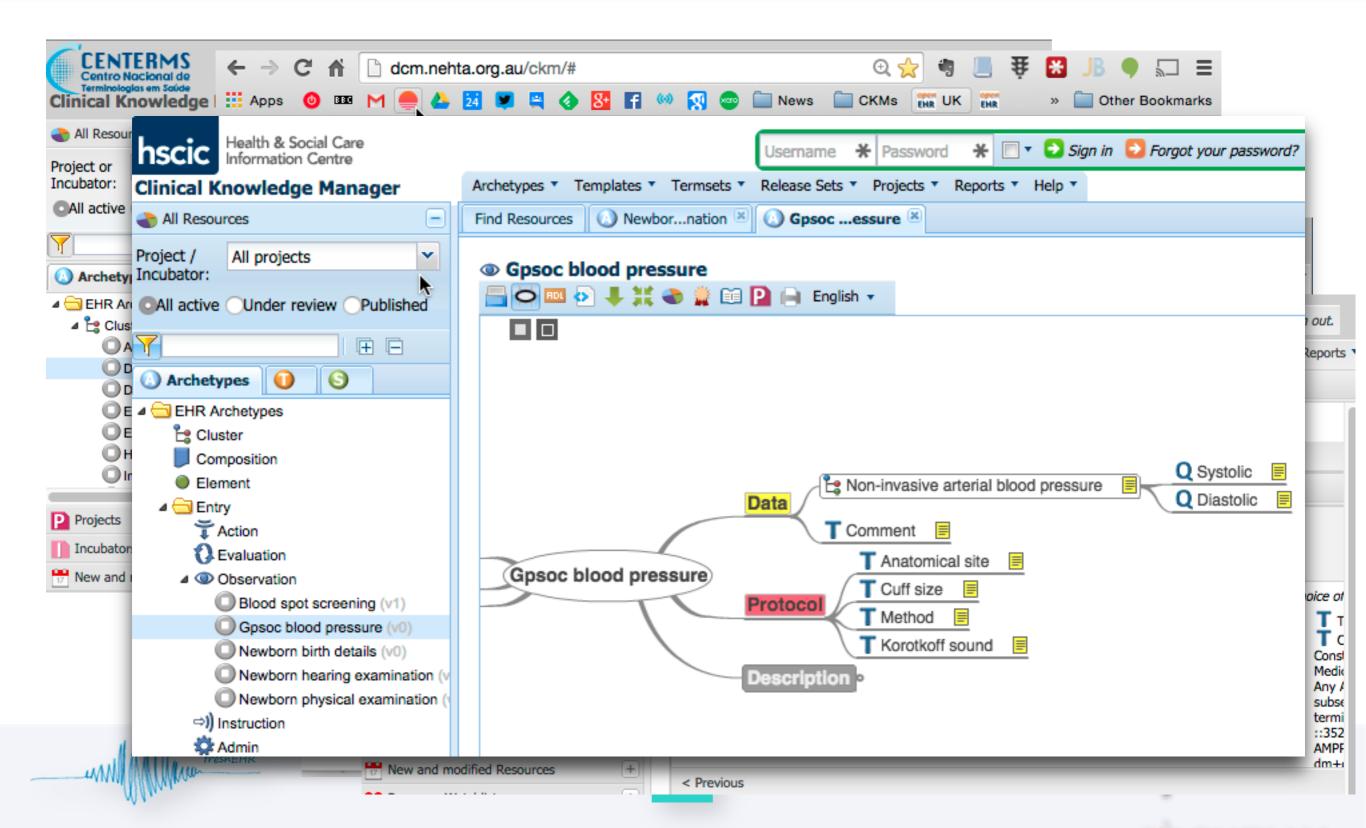
freshEHR







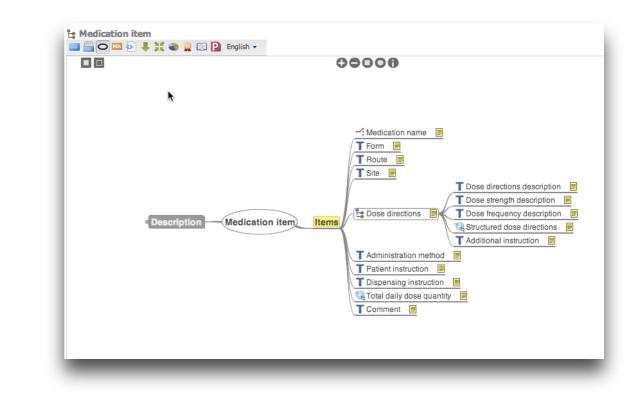
How is it used : National standards development



NHS Scotland

- Pioneered clinical-model based standards development
- Full commitment lacking
- Danger of losing 4-country solutions
- What is the alternative?

freshEHR





- MMMMMM freshEHR



