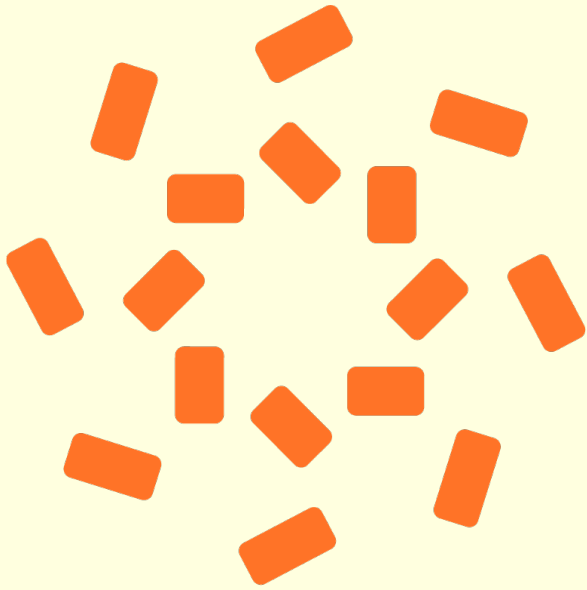


Pre meeting slide pack



Sparkled



Sparkled Clinical Content Design Group

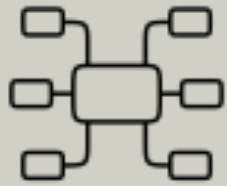
Tuesday 5 December Meeting

Online

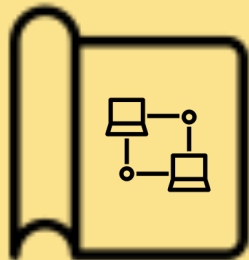


What is AU Core and Australian Core Data set for Interoperability (AUCDI)?

CDG is here



Specifies “*WHAT*” clinical information (and corresponding data elements and terms) should be included when sharing information supporting patient care



Specifies “*HOW*” the core set of data (above) and information should be structured, accessed and shared between systems

TDG is here

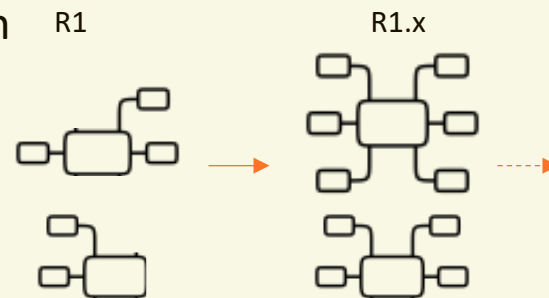


Sparked: AUCDI

Key outcome is “core of the core”

Start small and grow iteratively

- No simple “undo” – impact of change can be high
- Data elements can be added to over time
 - Work through backlog
 - Add more use cases
 - More functionality available



Stick with our design principles

Align and leverage work internationally were possible

- Where we differ, need to understand the impact



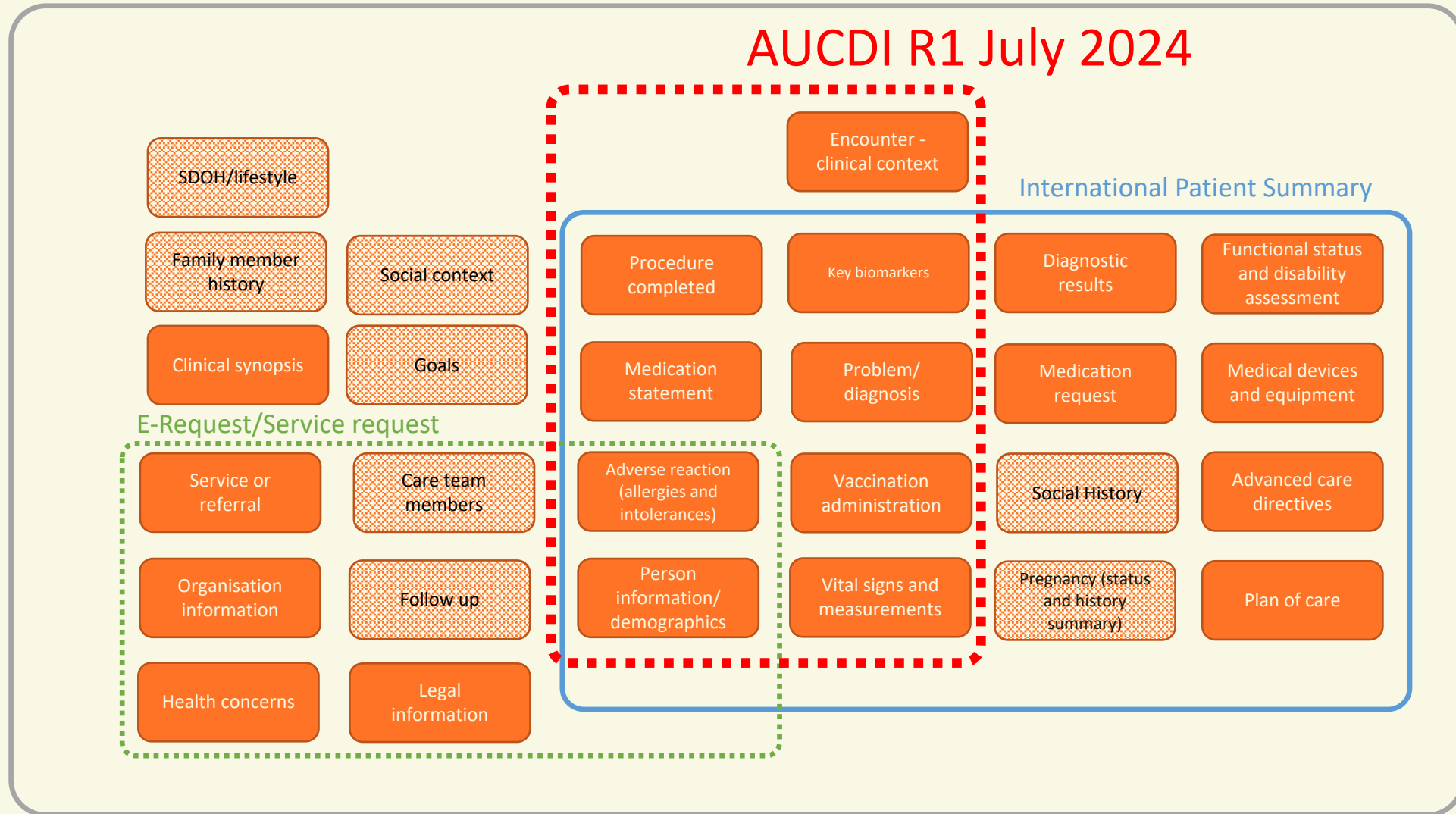
“From little things, big things grow”



Core Draft Principles of Data Set Design

- 1 Single entry, single development - multiple use and reuse
- 2 Supports patient centred care - driven by a clinical quality and safety use case
- 3 Reduce duplication of effort, not collecting of data for data's sake
- 4 Driven by primary clinical data use not secondary data use needs
- 5 Supports best practice care, clinical guidelines and clinician workflow
- 6 Systems can support now or with minimal effort, supporting a strategic roadmap with an agile iterative process
- 7 Leverage agreed national health data standards
- 8 Involve and consider all healthcare modalities

AUCDI – Updated Draft Scope (Dec 2023)





Proposed changes to scope of R1

- Clinical synopsis moved to “To Do List”
- Pregnancy status moved to “To Do List”
- Observations renamed to “Key Biomarkers” – scope limited to cardiovascular risk calculator for R1
 - HDL
 - LDL
 - Total Cholesterol
 - Trigs
 - HbA1c
 - eGFR
 - uACR



Proposed Candidate for Draft R1

Problem/Diagnosis

- Problem/diagnosis name
- Body site/laterality
- Status
- Comment

Adverse reaction risk

- Substance name
- Manifestation/s
- Comment

Gender

- Sex assigned at birth
- Gender identity
- Pronouns

Vital signs

- Blood pressure
 - Systolic
 - Diastolic
- Pulse
 - Rate
 - Rhythm*
- Body temperature
- Respiration
 - Rate

Procedure completed

- Procedure name
- Body site/laterality
- Clinical indication
- Date performed
- Comment

Medication statement

- Medicine name
- Strength
- Form
- Dose
- Route of administration
- Dose frequency
- Duration
- Instructions
- Clinical Indication
- Comment
- Status
- Start date
- Long term indicator

Tobacco smoking

- Overall Status

Measurements

- Height/length
- Body weight
- Waist circumference

Vaccination administered

- Vaccine name
- Sequence number
- Date of Administration
- Comment

Key biomarkers

- HDL
- LDL
- Total Cholesterol
- Trigs
- HbA1c
- eGFR
- uACR

Encounter – clinical context

- Reason for encounter
- Type of encounter



AUCDI

- What level of information should we include?

UK Professional Records Standards Body (PRSB)

Diabetes Record Information Standard
Version 1.0

['How to' Guide](#) Search By Item Filter By [Reset all](#)

MRO ⓘ	Name	Description	Value Sets ⓘ	Implementation Guidance ⓘ	Information Type ⓘ	Cardinality ⓘ
M	▶ Person demographics	The person's details and contact information.	-	This section contains the person's... Read more >	Record	1 ... 1
M	▶ GP practice	Details of the person's GP practice.	-	This section contains details of t... Read more >		1 ... 1

US Core Data set for Interoperability (USCDI)

DATA CLASS

Allergies and Intolerances

Represents harmful or undesirable physiological responses associated with exposure to a substance.

DATA ELEMENT	APPLICABLE STANDARD(S)
Substance (Medication)	<ul style="list-style-type: none"> RxNorm, January 6, 2020 Full Release Update
Substance (Drug Class)	<ul style="list-style-type: none"> SNOMED International, Systematized Nomenclature of Medicine Clinical Terms (SNOMED CT®) U.S. Edition, September 2019 Release
Reaction	<ul style="list-style-type: none"> SNOMED International, Systematized Nomenclature of Medicine Clinical Terms (SNOMED CT®) U.S. Edition, September 2019 Release

AU Primary Care Quality Design Foundations – Practice to Practice Use case

Data elements		
Substance	Description	<p>Identification of a substance, or substance class, which is considered to put the individual at risk of an adverse reaction event.</p> <p><i>Both an individual substance and a substance class are valid entries in 'Substance'. A substance may be a compound of simpler substances, for example a medicinal product.</i></p>
	Element occurrence	Mandatory, single occurrence
	Data type	<u>Codeable concept</u>
	Value set	This ADHA value set contains a broad set to support the recording of agents that may be responsible for causing adverse reactions. This set includes all AMT product concepts as well as SNOMED CT-AU substances such as chemical, material and dietary substances
	Examples	111088007 Latex 256259004 Pollen 102263004 Eggs (edible) 256349002 Peanut 373270004 Penicillin 3738011000036101 Amoxicil
	Alias	Agent, Class, Product

AUCDI

- Publish a logical data model
 - Similar to PCDQ
 - More than USCDI, closer to PRSB
 - Recommend terminology value sets where appropriate
- Will include
 - Name
 - Description
 - Element occurrence
 - Data type
 - Recommended standard code systems and value sets*
 - Alias*
 - Considerations*
 - Examples*

* Where appropriate or necessary

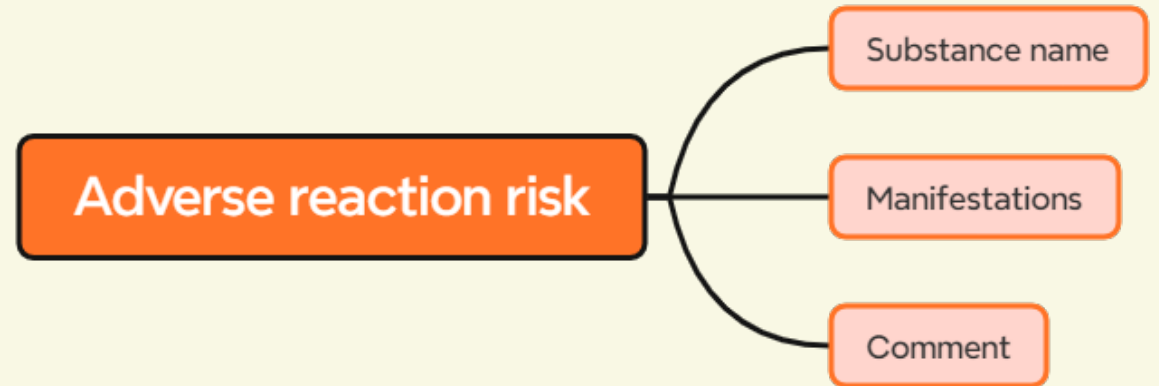
Into the data model...



Adverse reaction risk

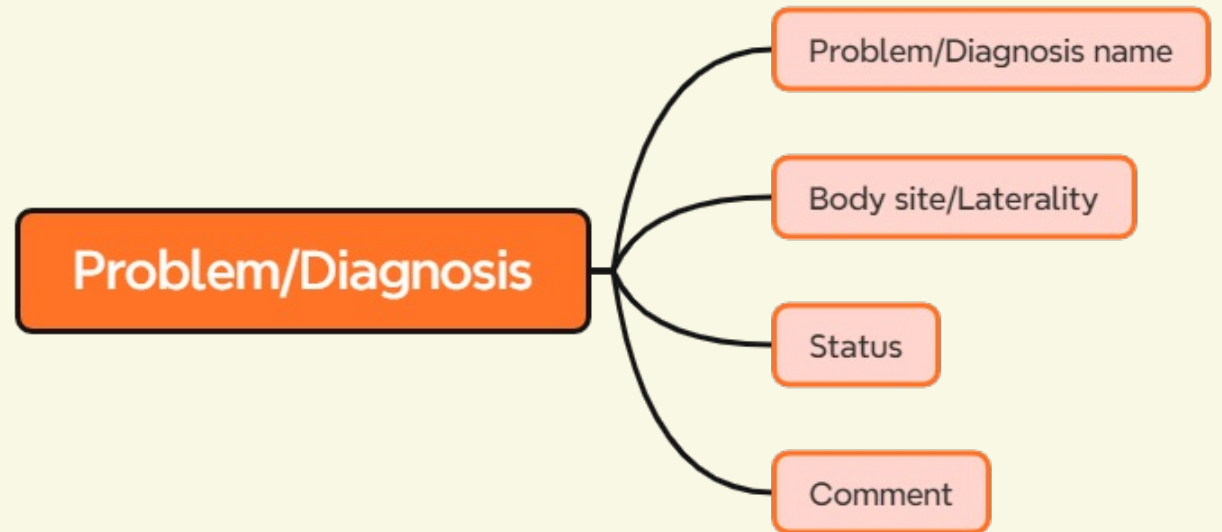
Details about a single substance (or class of substance) which, on exposure, may put the individual at risk of an adverse reaction.

- Includes, but not limited to, allergy and intolerance
- Scope:
 - Medication
 - Food
 - Environmental, including latex and bites/stings



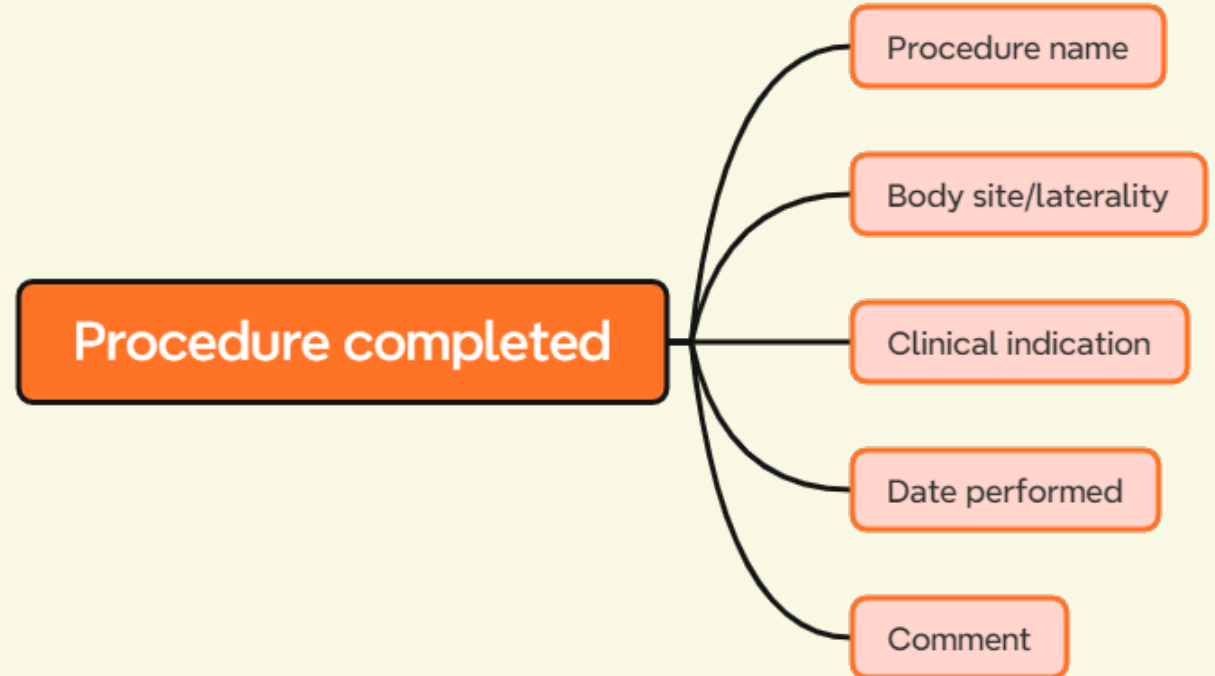
Problem/Diagnosis

Details about a single identified health condition, injury, disability or any other issue which impacts on the physical, mental and/or social well-being of an individual.



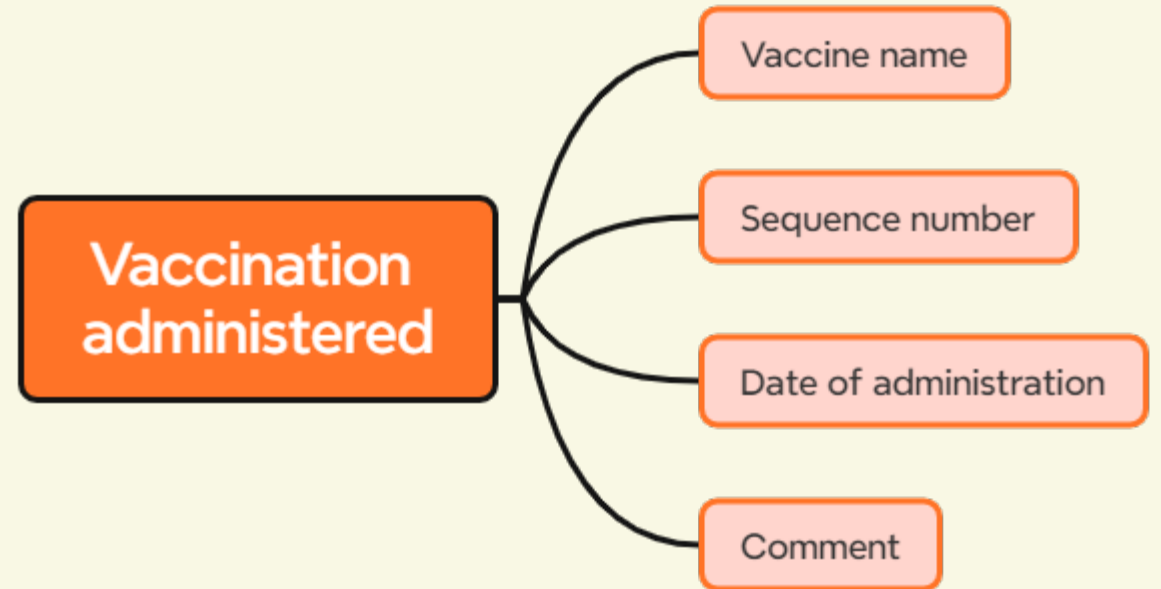
Procedure completed

Details about a single clinical procedure carried out for screening, investigative, diagnostic, curative, therapeutic, evaluative or palliative purposes.



Vaccination administered

Details about a single vaccine that has been administered to the individual.

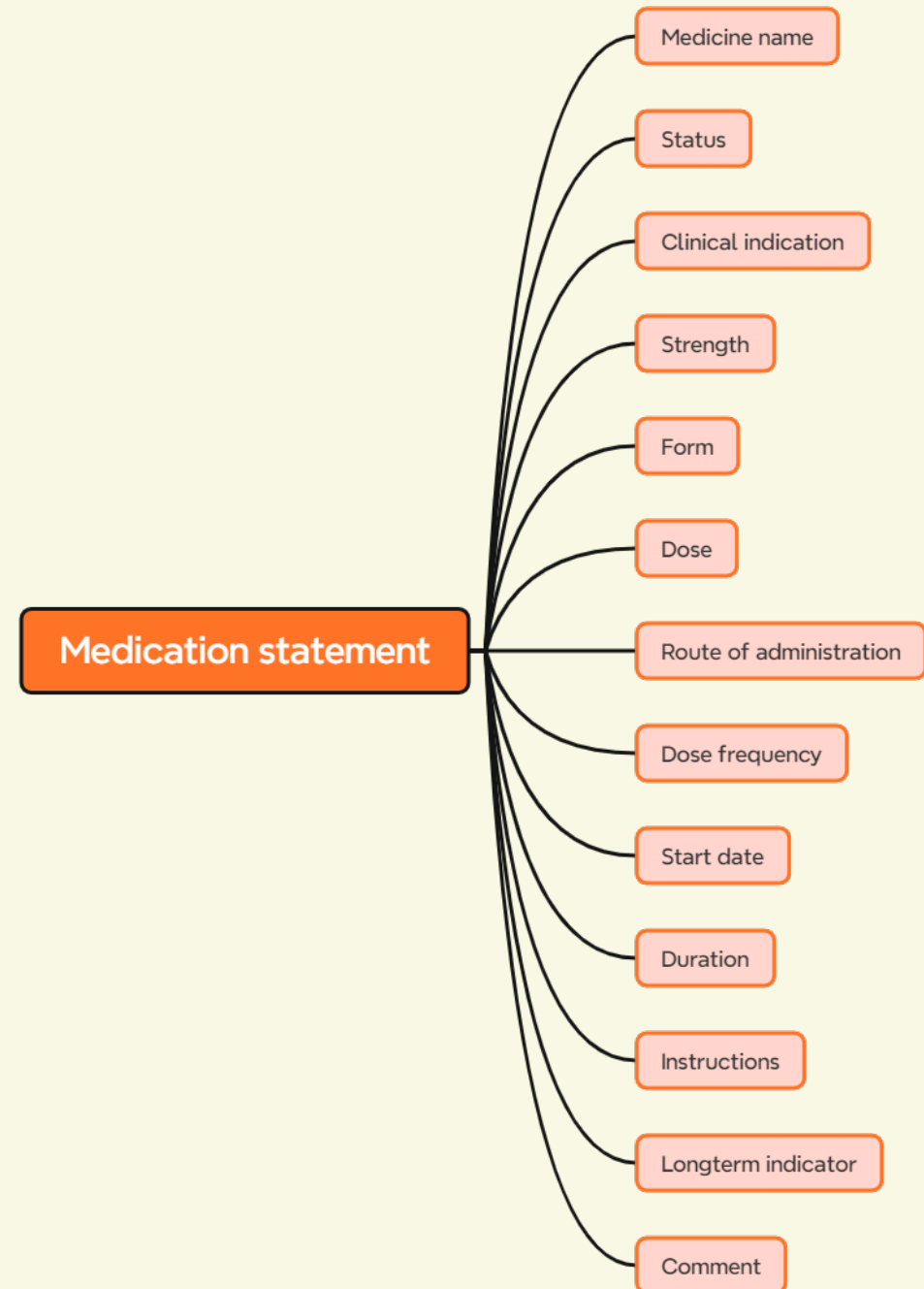




End of agenda for 5
December meeting
Next topics for 17
January meeting

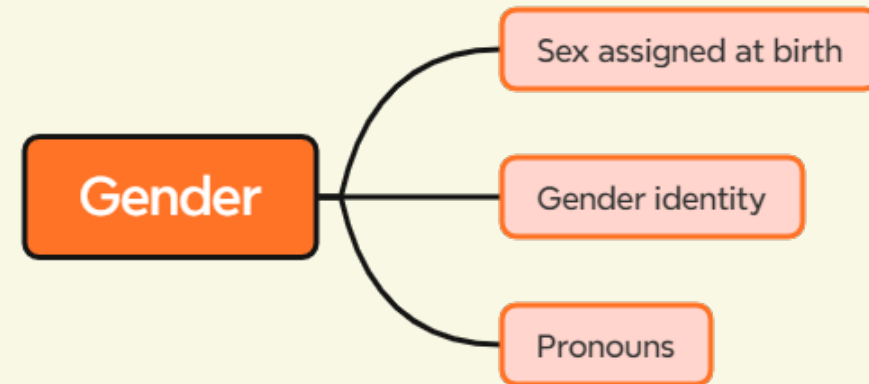
Medication statement

Details about a medicine, nutritional product or other therapeutic item that is currently being taken or used by the individual, including items that are prescribed or obtained over the counter.



Gender

- Sex assigned at birth
 - The sex of the individual determined by anatomical characteristics observed and registered at birth.
- Gender identity
 - The individual's internal perception of their own gender.



Tobacco smoking

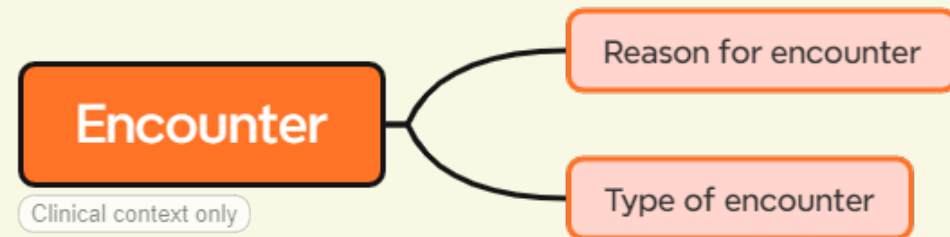
Details about the tobacco smoking habits of the individual.

- Statement about current smoking habits for all types of tobacco.
- Proposed value set: Never, Current, Former.

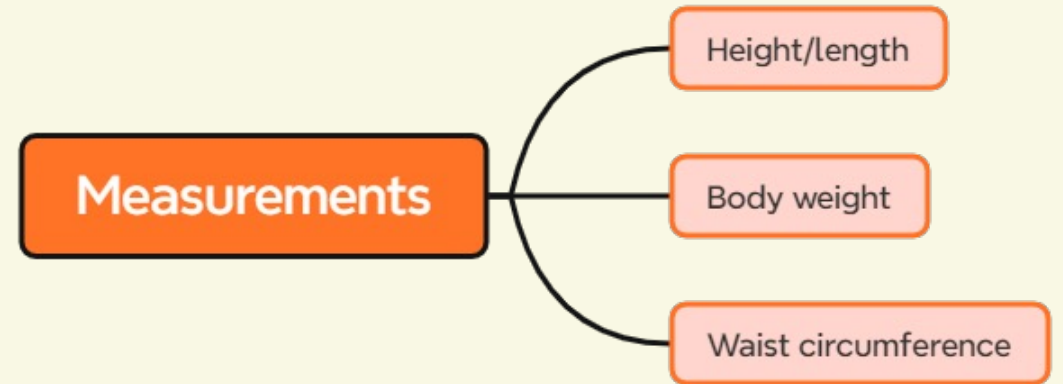
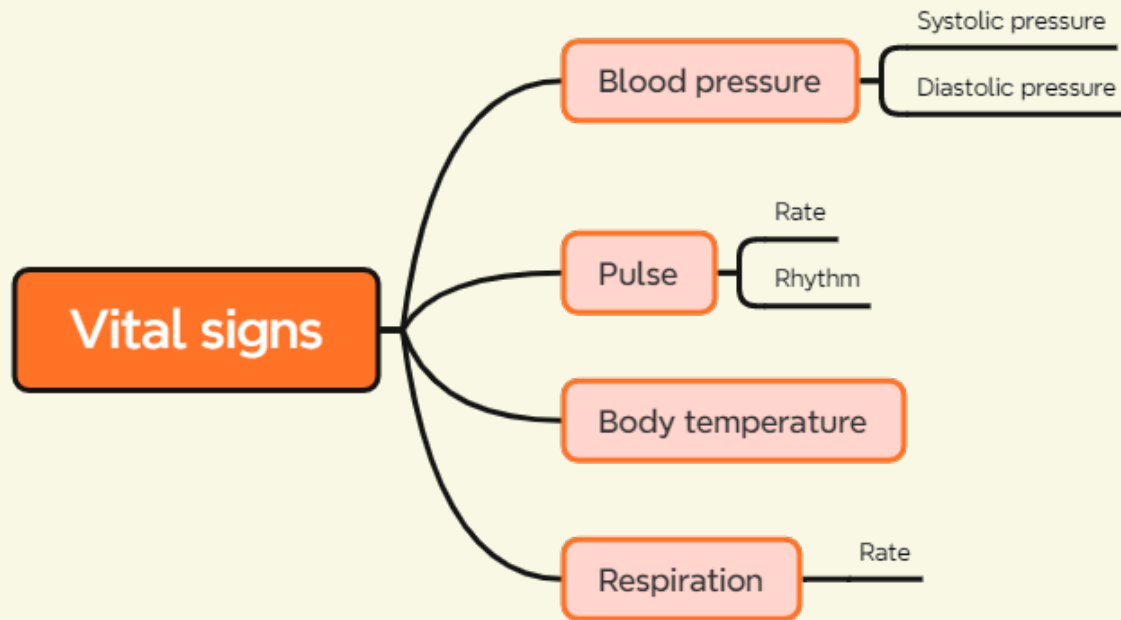


Encounter – clinical context

- Reason for encounter
 - The reason for initiation of any healthcare encounter or contact by the individual who is the individual.
 - Proposed value set combines clinical and administrative values
 - ‘Presenting complaint’ (or similar) with a value set comprising symptoms and signs, for example ‘shortness of breath’ or ‘vomiting’; and
 - ‘Reason for attendance (or similar) with a value set comprising terms that reflect administrative, billing or funding reasons, for example ‘pre-employment medical’ or ‘routine immunisation’.
- Type of encounter
 - The modality through which the encounter occurred or was conducted



Vital signs and Measurements



Key biomarkers

- Initial R1 scope to support calculation of cardiovascular risk assessment

