

Med-e-Tel 2006

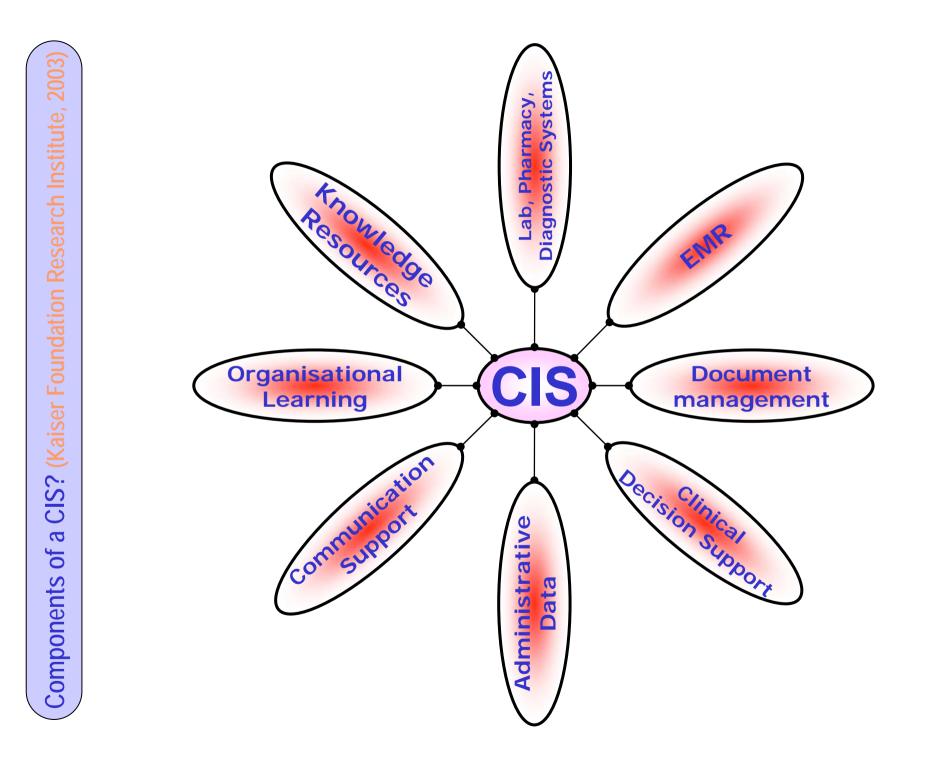
Luxemburg 05.04.2006

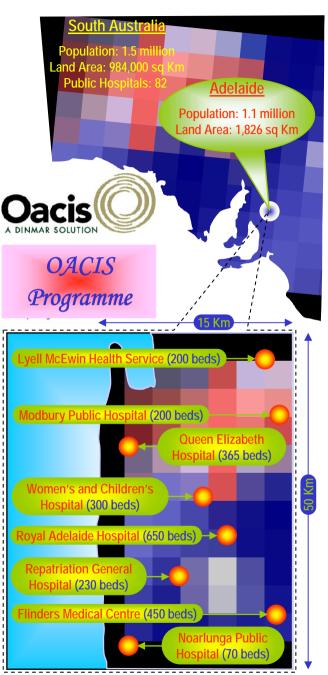
Progress of e-Health in Australia: OACIS

(The Open Architecture Clinical Information System)

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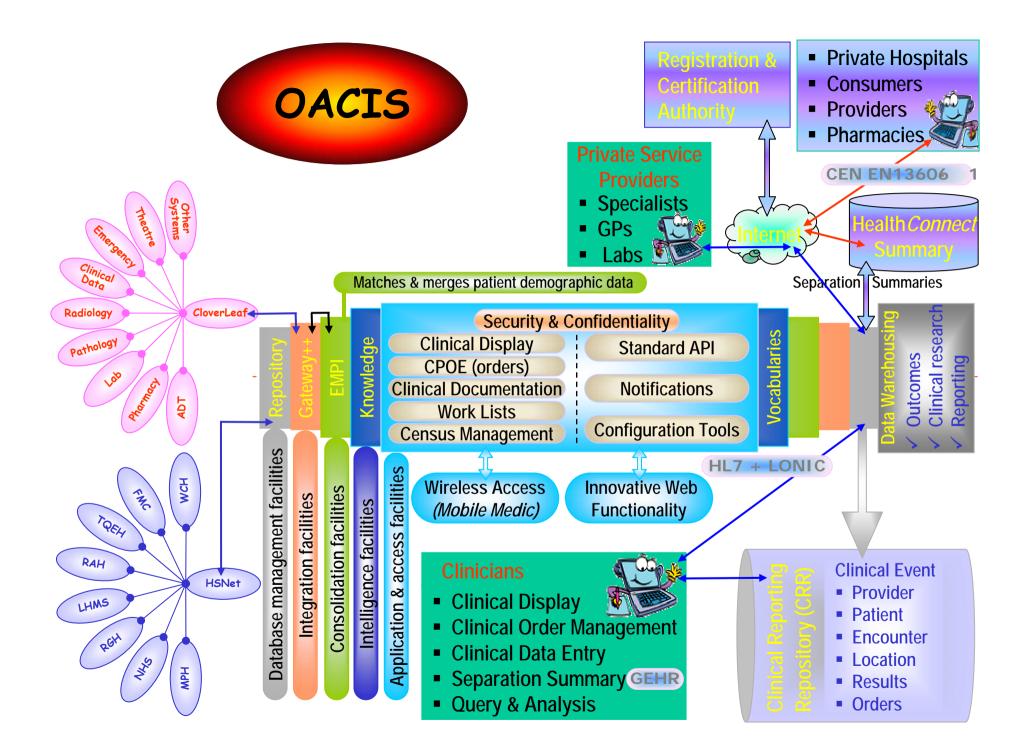
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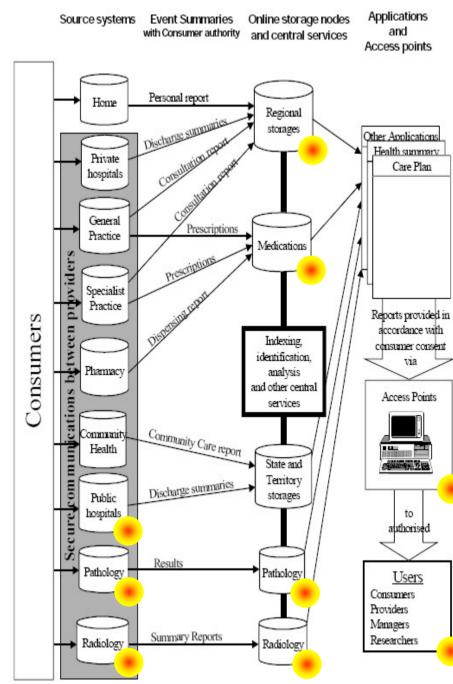




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	1990	DHS identifies the need to improve CIS, to enable clinical best practices and thereby improve quality of care.
n m	1995	The Renal Information System at TQEH required replacement.
		A world wide study was conducted looking at both clinical and technical requirements.
		DHS selected OACIS out of west cost of the USA, now owned by DINMAR in Canada.
	4007	
	1997	Pilot implementation by the Department of Health's ICT Services <i>Projects Branch</i> in partnership with Accenture in the renal unit of 4 (TQEH, RAH, FMC, WCH) of the metropolitan public hospitals to test and evaluate benefits and outcomes able to be achieved.
	1999	OACIS Enterprise Wide Steering Committee established.
	2001	Implementation of data conversion between OACIS to GEHR and GEHR to OACIS.
		Extended implementation of OACIS across 8(TQEH, RAH, FMC, WCH, LMHS, MPH, RGH, NHS) major metropolitan public hospitals.
	2004	Participation of SA in the implementation of first stage of Health <i>Connect</i> .





DHS is represented on the Health Connect Board

DHS and SADI (South Australian Division of General Practice Inc.) have jointly developed a state-wide GP Register

Health Connect's Clinical Information Project:

- Developing a framework for defining clinical information capture (in the form of 'event summaries' and representation (lists, views and reports) in shared EHRs
- The initial health profile, GP consultation, hospital discharge summary, pathology, and imaging event summaries were identified as immediate priorities

OACIS to GEHR and vice versa done by GPCG.

Stakeholder participation:

 Clinical Reporting Repository Committee with representation from DHS, hospitals, academia, GPs, consumers and legal

Use of electronic signatures for drug prescribing and also procurement

Ensuring Privacy/Confidentiality by adopting the Code of Fair Information Practice

The Department of Health of South Australia has adopted a Code of Fair Information Practice to ensure that all public hospitals and health units comply with a set of Privacy Principles.

Collection, storage & transfer, usage and disclosure of personal health information are regulated by principles.

Providing personal health information is voluntary^{*} but staff have obligation to record details of service they provide.

Only required information is passed on to staff who are involved in your care.

Under some laws, such as the Public and Environmental Health Act, the South Australian Health Commission Act and the Motor Vehicles Act, doctors are required to report certain information.

Some authorities such as HIC is legally entitled to obtain certain information.

Except for some legal exceptions, under Freedom of Information Act, an individual can access his/her medical record to get a copy or to amend the same.

	OACIS Programme	HealthConnect
What?	Hospital based clinical information management for renal patients	National system of EHR to improve flow of information across the health sector
Focus	Better decision making in acute care	Providing valuable information for health sector
Propagators	Patient centric and provider/clinicians driven	Commonwealth and State and Territory governments driven
Scope	Limited to Adelaide metropolitan public hospitals + some GPs and Private units	Throughout Australia
Collection	Comprehensive encounter information from source systems	Brief event summaries from point of care
Storage	Held in central storage repository	Held at regional storage service close to point of care
Ownership/Funding	South Australian public health system	Jointly Australian, State and Territory governments
Architecture	Client-server under Intranet + Some have web access (separation summary)	Web based under secure network using PKI
openEHR	HL7 messaging + LOINC code sets	CEN EN 13606-1 (CEN archetypes or HL7 templates)
Consumer	 Consumer not registered Implied consent model used for paper based records Medical records are linked by matching patient demographic information Access to information under Freedom of Information act 	 Voluntary participation, Unique ID after registration Information storage/disclosure with individual's consent Event summaries and medication details are linked by using consumer ID Complete control and on-line access to information
Provider	 Unique logon ID for all providers involved in patient care GP registry prepared by SADI Providers to follow set of Information Privacy Principles based on Privacy Act 1988 Only authorized providers will password authentication (based on role and site) can access information Only required information connected with patient care is made available 	 Voluntary participation, Unique ID after registration National provider directory being prepared at WA Providers to follow 10 National Privacy Principles till National Health Privacy Code is introduced With patient's consent, only authorized providers can access information With patient's consent, only required information connected with patient care is made available