Conference papers

Regional repositories, reintermediation and the new GMS contract: cardiovascular disease in Tayside

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ABSTRACT

Background The new contract for general medical practitioners will make increasing demands on the primary care informatics community. There are a number of potential ways to provide reports which meet the requirements for data on the quality of care being provided by practices. In Scotland there are four components of the national information technology strategy which make meaningful comparisons of data possible.

Objective Using cardiovascular data as an example, to describe how the community health index number, managed clinical networks (MCNs), increasing consistency of Read codes, and regional repositories of data make the acquisition, processing and use of data more straightforward.

Method The cardiovascular MCN collects the majority of its data electronically and four properties are crucial to its success: automatic collection of electronic data from many sources, prioritisation of data derived from multiple sources, record linkage processes, and manual validation of electronic data.

Results Clinicians in primary and secondary care enter data during consultations and see the results of consultations recorded elsewhere. Because all data from the region are able to be Read coded according to prespecified templates, we are able to indicate to practices where they are in relation to the new contract targets and indicate which patients need to be seen, or excluded from, calculations.

Conclusion Effectively integrated management is facilitated by provision of regular prompted recall and review of people with chronic disease by multidisciplinary teams collaborating across the health service and into the community. In Scotland, use of newer informatics tools are proving to be useful contributions from primary care computing to equitable, evidence-based care.

Keywords: computerised medical records, coronary heart disease, data quality, general practice, quality improvement

Introduction

The recent acceptance of the new General Medical Services (GMS) Contract by United Kingdom general practitioners (UK GPs) presents challenges and opportunities to the primary care informatics community. All serious system vendors have promised to

provide users with computer systems that will meet the demands of this new contract for reports that meet the requirements of reporting systems.¹ These will satisfy the demands of individual practices, but there may be difficulties in making comparisons between different practices within the same primary care organisation (PCO), region or nationally if different data definitions are used.

In Scotland there have been four developments that make meaningful comparisons more straightforward: use of a unique patient identifier (community health index [CHI] number), managed clinical networks (MCNs), increasing consistency of Read codes, and regional repositories of data. Patients who register with a GP in Scotland are allocated a unique ten-digit identifier, known as the CHI number, which incorporates date of birth and sex.2 Access to the Community Health Master Patient Index enables patients to be tracked as they move between areas within Scotland. Because the CHI number is unique, it enables direct matching of index cases to subsequent events.3 The key concepts and core principles of MCNs in Scotland were set out in a management executive letter and have been described in greater detail elsewhere.^{4,5} They were defined as 'linked groups of health professionals and organisations from primary, secondary and tertiary care, working in a co-ordinated manner, unconstrained by existing professional and health board boundaries, to ensure equitable provision of high-quality, clinically effective services throughout Scotland'. In Scotland, Scottish Clinical Information Management in Primary Care (SCIMP) was set up in 1999 in order to provide a central body to co-ordinate and standardise the management of clinical information in primary care.6 The aims of the group were to publicise the benefits of a co-ordinated Scottish approach to data management using Read codes.⁷ Regional repositories of data have been a key principle of the Scottish information management and technology (IM&T) strategy for the past five years.8 It is the intention of the National Health Service (NHS) Scotland to create regional repositories of data that enable integrated care throughout the country.9

In the Tayside region of Scotland, incorporation of the CHI number within routinely collected primary and secondary care data is almost 100%. The HEARTS (Heart Disease Evidence-based Audit and Research Tayside Scotland) MCN has extended the already successful DARTS (Diabetes Audit and Research Tayside Scotland) model of data sharing between primary and secondary care to coronary heart disease (CHD).10 In diabetes this model has already become the national system, known as SCI-DC.11 Originally configured as a secondary prevention project, HEARTS MCN now extends to other forms of CHD.¹² This paper describes the process of reintermediation whereby data is collected, validated, prioritised, linked and returned to practices as more useful information for the practice, PCO and region.¹³

Methods

HEARTS MCN collects the majority of its data electronically and four properties are crucial to its success:

- the ability to automatically collect electronic data from many sources
- prioritisation of data derived from multiple sources
- the ability to operate and link patient records regardless of the unique patient identifier (UPI) scheme(s) in place, i.e. the CHI number
- manual validation of electronic data.

Automatic data collection

Automating the collection of information from different health computer systems in the NHS, and elsewhere, remains difficult.¹⁴ The HEARTS MCN addresses this problem through the use of a tool known as GENIE (Generic Importer and Exporter), developed with DARTS, which greatly simplifies the task.¹⁵ GENIE manages the overnight transfer of new or modified data from the practice to a regional computer, where the files are uncompressed and unencrypted. HEARTS MCN takes these data and merges them with patient information already held on the database.

Prioritisation of data from multiple sources

Through a process of consultation between the professions in primary and secondary care, the quality of each data source is determined. Each data source is then allocated values indicating its various abilities to provide information on different aspects of disease. When displaying or summarising data, HEARTS MCN uses the best and most relevant source of information at all times. For example, the cardiology clinic is more likely to generate accurate information regarding risk stratification than a general practice. Conversely, a practice is more likely to be accurate regarding the current treatment regime. Cardiology clinic records then take precedence where there is any ambiguity regarding the state of a patient's risk status, and a general practice record would take precedence when describing a patient's current treatment regime.

Patient identifier linkage

To operate across political boundaries, a system must be capable of dealing with multiple patient identification schemes and of dealing with patients who have more than one number even within a consistent numbering scheme. The system we use supports any number of patient identification schemes and there is no limit on the number of identifiers any one patient can possess.

Manual data collection and validation

Since comprehensive electronic records are rare and we want to provide a high quality of data to support equitable care, some data must be collected manually. HEARTS MCN minimises this task by identifying data collection holes for a facilitator to target. The facilitators have several roles, including a responsibility for disseminating patient/clinic/practice information, providing a human interface to the project, identifying training needs and aiding in general facilitation.

Results

Clinicians in primary and secondary care are able to enter data during consultations (see Figure 1), and see the results of consultations recorded elsewhere (see Figures 2 and 3).

Because all data from the region are Read-coded according to pre-specified templates, we are able to indicate to practices where they are in relation to the new contract targets (see Figure 4), and indicate which patients need to be seen or excluded by drilling down into the list of those for whom the target level has not been reached (see Figure 5).

Discussion

The new GMS contract will reward practices for their performance against a set of 76 indicators covering ten disease groups, including CHD. Between 30% and 50% of GPs' income may be dependent on meeting the targets specified in the indicators. The informatics tools developed in Scotland and Tayside over the past five years have enabled rapid access to data on quality of care for CHD for the new GMS contract. This has allowed us to develop a system that allows data sharing for a variety of approved purposes. The UK Data

NHS Tayside	Tay Bridge Medical Centre (11058) HEARTS Clinical Review Form					
Patient / Practice Details Review Date Patient ID Annual Review	24 / 07 / 2003 260342UOMR	Hospital ID Administration	e Health Board Number Form	11058 CHI		
Forename Clinical History Most Recent MI 18 / 10 / 1999 Date Hypertension Diagnos / / / Date LV Dysfunction Diagn		Most Recent Coronary Angioplasty 29	Most Recent CABG // // Type of Diabetes Not Diabetic Date Diabetes Diagn	BELL		
	/ (mmHg) III	BMI Lab Investigations Blood Glucose Fasting Blood Glucose Total Cholesterol Ankle Oedem	5.4 mmoVL (19/02/2002) IIII No result stored 4.9 mmoVL (19/02/2002) IIII a? CYes CNo IIII			
CLASP Score Short of Breath? CLASP Score Lifestyle Alcohol Intake (Units P/W) Advice Given Diet Advice Given	Smoking (Current See Ex Smok	Smoker ser pped 01 / 01 / 1987 noked	Exercise (MG) C Physically Impossibl C Avoids Exercise C Light Exercise C Moderate Exercise C Heavy Exercise	le		

Figure 1 HEARTS clinical review form



Home: Search: Patient Search Results: Echo results for an individual patient: Condensed ECHO Report

This is the condensed version of the ECHO report;

Link To Complete Version of ECHO Report

Name:	Patient Identifier:	Ward-line Cardiac Op	Respital Name: Perth Royal Infirmary	Date of Echec 23/04/2003
Consultant :	Dr P Currie		Reason for Request :	LV+Valve assessment
Aortic Valve Appearance :			Aortic Regurg :	
Aortic Stenosis :	No stenosis			
Mitral Valve Appearance :			Mitral Regurg :	
Mitral Valve Comments :	Reduced motion of Ant MVL due	to eccentric MR jet		
Fricuspid Valve :	Normal		Tricuspid Valve Regurg:	
Pulmanary Valve :			Estimated PA pressure:	0 mmHg
eft Ventricle:	Normal systolic function			
Summary Information :	Left atrium normal		RA/RV normal	
Overall Comments :	Echo & report reviewed by Dr Cu	rrie Significant aortic regurg	tation. Unchanged from previous exam.	

1 Record(s) Found.

Website Comments

WY WAI-AAA WY COS.

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Figure 2 Condensed echocardiogram (ECHO) report



Home: Practice Overview: Patient Population: Patient Summary Screen: ETT Results for an Individual Patient: Complete ETT Report

This is the Complete version of the ETT report;

Patient Name:	BROWN, PHILIP	Patient CHE	2109470WWH - CH	GP Name:	Dr BERENHOLLINS	Consultant:		Johnstone, Dr D
Wd / Clinic :		5/6 OP		Hospit	al:	Ninewells		
Record Date :		10/03/1998						
Current Cardiac	Medication (One):		To	est Indication :	Diagno	sis of chest pain		
Two:			Test Type : Full		Full Bi	uce Protocol		
Three:		To	Technician :		5			
Max ST Elevation	n Resting :				0 mm	Lead:		
Max ST Elevation	n on Exercise:				0 mm	Lead:		
Max ST Depr Re	sting :				0 mm	Lead:		
Max ST Depron	Exercise :				0 mm	Lead:		
ST Depresion :								
Resting HR:		66 / min	Rest	ing BP :	145/8	5 mmHG		
Standing HR :		77 / min	Max	BP:	160/9	0 mmHG		
Max HR :		141 / min						
Pred. Max HR:		169 / min			% MaxHR:		83	%
Exercise Time :	xercise Time: 4 mins 39 s		ecs	METS:			6	
Comments:								
Reported By :				Dr	Sim			

Figure 3 Complete exercise tolerance test (ETT) report

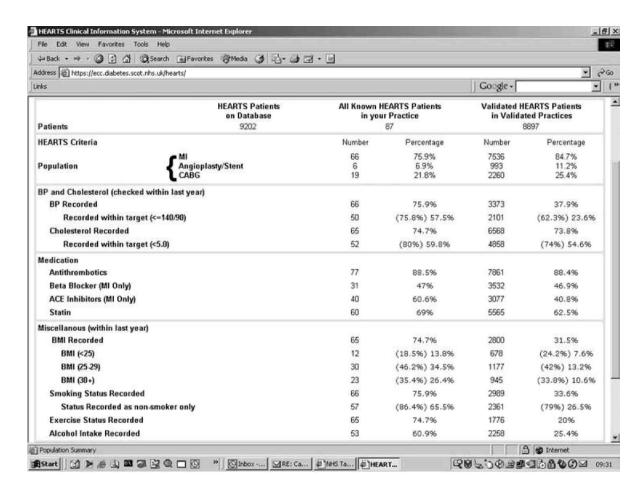


Figure 4 GP quality indicators summary page

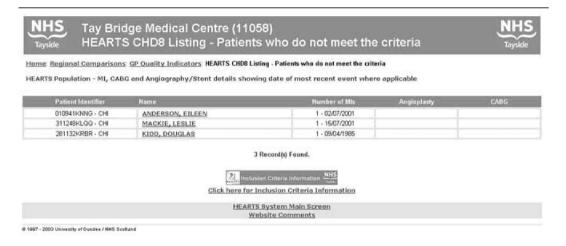


Figure 5 Sample drill down for CHD

Protection Act of 1998 has led to major difficulties for everyone engaged in record linkage of patient data. 16,17 HEARTS MCN employs a variety of approved mechanisms to maintain security and the rights of patients.

- All communications involving confidential data are encrypted.
- Firewalls and subnet address restrictions help prevent inappropriate access via TCP/IP (Transfer Control Protocol/Internet Protocol) and the Internet.
- All attempts to log-on to HEARTS MCN and every action subsequently taken are logged, producing audit trails.
- Patients must be centrally registered as belonging to a practice before that practice can view that patient's data.

The patient is ultimately in control of who can access their records. HEARTS MCN supports informed consent though the use of consent granting and denial forms. Should consent be denied, all records pertaining to that patient, with the exception of that patient's unique identifiers, are deleted. Once consent has been denied, that individual's care is conducted using only their written records.

We use an approved cleaning and anonymisation process (CLAM) to de-identify data items such as the CHI, references to a GP either by the NHS identifier or the General Medical Council (GMC) registration, references to a GP practice and references to a pharmacy. With these references anonymised, very nearly all the remaining data that could immediately identify individuals can simply be removed from the datasets. For example, all names and addresses and hospital or practice names can be removed without losing any information that might be of use to researchers.

The exact role that quality indicators in the new contract will play remains contentious. 19,20 While GPs are likely to respond to financial incentives aimed at improving quality of care, there remains an important question as to how much total health gain is likely to result from the measures chosen in the new contract.²¹ Use of quality indicators in other countries has sometimes led to perverse incentives or gaming.²² What appears to be required for effectively integrated management of chronic diseases such as CHD, however, is provision of regular prompted recall and review of people with chronic disease by multidisciplinary teams collaborating across the health service and into the community. In Scotland, use of a UPI, the emergence of MCNs, more consistent recording of Read codes and regional repositories of data are proving to be useful contributions from primary care computing to equitable, evidence-based care.²³

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CONFLICTS OF INTEREST

None.

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