RESEARCH



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The use of a global trigger tool to inform quality and safety in Australian general practice: a pilot study

Background

Systems to identify risks and adverse events (AEs) in Australia are limited. This study aims to explore whether general practice records contain information on AEs, and to conduct a pilot study on the type and frequency of AEs in general practice in Australia, using a global trigger tool (GTT).

Methods

Five practices were recruited and consented to collect data. Practice nurses were trained to collect data at their practices. Records from randomly sampled patients aged 75 years or older were reviewed.

Results

A total of 428 patient records were reviewed. A total of 44 AEs were detected in 41 records. The percentage of patients with an AE was 9.6%. Most low preventability AEs (21/29) were medication incidents.

Discussion

The study found that significant levels of information about AEs exist in general practice medical records and rates of harm are broadly in line with a similar study in Scotland.

Keywords

quality assurance, health care; risk management; general practice

General practice provides 116 million consultations to Australians every year at an average of 5.3 visits per head of population.¹ However, evidence for safety priorities and systems to identify, analyse, respond to, monitor and prevent risks and adverse events (AEs) in Australian general practice is limited.^{2,3} Providing data on harm may be an enabler to engage general practitioners (GPs) in safety and quality activities such as quality improvement projects.

A retrospective review of medical records can provide data on the frequency and type of AEs.⁴ One method, known as the global trigger tool (GTT), initially developed by the Institute of Healthcare Improvement (IHI) in 2003 for use in hospitals,⁵ has been modified for primary care in the United Kingdom^{6–9} and North America.¹⁰ There are no published studies using a GTT in an Australian general practice environment.

GTTs use a series of triggers to screen the medical record for a potential AE. The presence of a trigger then leads to a more in-depth review of the record. AEs are then coded according to their type and an incident rate is calculated. The original IHI version of the GTT has been shown to have high specificity, moderate sensitivity and favourable inter-rater and intra-rater reliability using hospital-based records.¹¹

GTTs can be used to identify and prioritise types of AEs, such as medication errors, which may subsequently be addressed with quality improvement processes. Data collected at one practice can potentially be aggregated to regional and national levels.¹²

The aim of this study was to explore whether general practice records contain

information on AEs, and to conduct a pilot study on the type and frequency of AEs in general practice in Australia, using a GTT.

Methods Definitions

Definitions for a patient safety incident, AEs and harm were derived from the International Classification for Patient Safety (ICPS).¹³

Triggers

A two-stage Delphi process was used to develop the triggers, using GPs and practice nurses who were members of a local Division of General Practice Clinical Leadership Group (CLG) as experts. Twenty-eight candidate triggers, modified from primary care GTTs developed in Scotland⁶ and England,⁸ were the starting point and a final list of ten triggers (*Table 1*) was produced.

Development of a data collection tool

A paper-based data collection tool was then developed, which recorded the presence of triggers and whether an AE occurred for each record review. For those record reviews with an AE, preventability (low, medium, high), level of harm¹⁴ and AE source (primary or secondary care) were recorded. A free-text field to describe the type of AE was also included. The patient's gender and frequency of general practice service use were collected for denominator purposes. A user guide outlining definitions, sampling method and data collection methods was also developed.

Recruiting

Five practices were recruited from within the Southern Adelaide – Fleurieu – Kangaroo Island

Medicare Local (SAFKIML) region and each consented to collect data on 100 patients each. The only inclusion criterion was that practices used electronic clinical software. Each practice nominated a practice nurse to undertake the reviews and one researcher (PH) trained the nurses in the method.

Sampling and reviewing

Records were sampled from patients aged 75 years or older as the CLG provided advice that this group may be more vulnerable to harm if exposed to a patient safety incident. Records were reviewed from randomly sampled patients who had attended the practice three or more times in a 6-month period (January–July 2012).

Patient records were reviewed for the presence of triggers and, subsequently, for AEs if a trigger was present. No patient identifiers were recorded. Data collection forms were then securely mailed to the research office where they were entered into a Microsoft Access database and analysed.

Ethics

Ethics approval was sought and received from the Royal Australian College of General Practitioners (RACGP) National Research and Evaluation Ethics Committee (NREEC 12-001).

Results

Records of 428 patient (225 females and 203 males) were reviewed. There was a total of

Table 1. Final list of 10 triggers					
Trigger	Trigger grouping				
Were there three contacts with the practice in any given period of a week (this includes telephone calls, consultations with nurse/GP or home visits)?	Consultations and attendance				
Had the patient been admitted to a hospital for any intervention, management or procedure (including elective surgery) for at least one night?	Consultations and attendance				
Did the patient have a consultation out-of-hours or attend emergency or an ACIS callout/attendance?	Consultations and attendance				
Was the patient prescribed opioid analgesia?	Medication				
Was the patient prescribed a benzodiazepine?	Medication				
Was the patient prescribed an antipsychotic?	Medication				
Was the patient prescribed NSAID/COX ₂ inhibitors?	Medication				
Were there any INR readings >4.5?	Laboratory				
Was the eGFR $\leq 60 \text{ mL/min}/1.73 \text{ m}^2$?	Laboratory				
Was the haemoglobin level ≤100 g/L?	Laboratory				
ACIS. Acute Crisis Intervention Service: COX ₂ , cyclooxygenase 2; eGFR, estimated					

ACIS, Acute Crisis Intervention Service; COX₂, cyclooxygenase 2; eGFR, estimated glomerular filtration rate; INR, international normalised ratio; NSAID, non-steroidal antiinflammatory drug

Table 2. Number of medical record reviews by practice					
Practice number	Number of medical record reviews	Reason for practice not completing 100 records			
1	86	Insufficient number of eligible patients			
2	98	Two missing forms			
3	100				
4	100				
5	44	Insufficient number of eligible patients			
Total	428				

4117 patient consultations (9.6 per patient), the majority (73%) being in-room consultations with the GP. Three practices did not complete 100 records because of either an insufficient number of eligible patients or missing forms (*Table 2*). The average time taken to review a medical record was 4.5 minutes (*Figure 1*).

Triggers and AEs

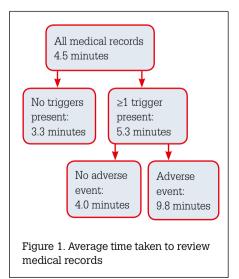
Table 3 shows the number of AEs by triggers and medical records reviewed. 273 medical records (64%) were coded as having one or more positive triggers (range 1–7) and of these 44 AEs were detected in 41 records (*Table 4*).

The percentage of patients with an AE was 9.6%. Primary care was the point of origin in 26 AEs (59%), and 18 AEs (31%) occurred in other levels of care. Reviewers deemed four AEs (10%) as highly preventable, nine (21%) moderately preventable and 29 (69%) low (with two not scored). Most low preventability AEs (21/29, 72%) were medication incidents, usually with unknown adverse reactions.

Discussion

Main findings

Using a consensus-based approach, this study modified an existing tool for an Australian general practice environment to develop a set of triggers designed to detect AEs. Trained staff then reviewed records for triggers and AEs. The method found an AE rate of 9.6 per 100 patients over a 6-month period;



66% of AEs were medication-related. This is consistent with a similar study in Scotland that found a rate of 9.4% with 59% being related to medication.⁶ Sampling strategies in this study (patient age >75 years) were different from the study in Scotland⁶ (patients age >18 years). Another study in primary care in North America found medication-related AE rates of 15%.¹⁰

Balancing sensitivity and feasibility

Although retrospective review of medical records is recognised as one of the more sensitive methods of collecting AEs, it can be resource-intensive and time-consuming.⁴ This is a major barrier to GPs or organisations undertaking their own reviews.

Only five triggers had a positive predictive value of more than 15% and two of these (haemoglobin and antipsychotic medications) were associated with low numbers of AEs. Those triggers that occurred frequently and had a high positive predictive value (such as

three general practice contacts in one week, a hospital admission and an after-hours consultation) yielded higher numbers of AEs and require further research. Another avenue for further research into a more time-efficient method may be the use of electronically detected triggers. Medication-related AEs^{9,10,15} and diagnostic errors¹⁶ have been detected using electronic methods.

Policy implications

A national discussion paper entitled *Patient Safety in Primary Health Care*² found that one of the four key themes emerging from a consultation process was a lack of knowledge and understanding of patient safety risks in primary healthcare, and limited coordinated systems to identify, analyse and respond to AEs. The paper proposed 'developing a systematic and coordinated identification, reporting and monitoring system for patient safety incidents in primary care. It was suggested that any activity in this area should include infrastructure and resources to implement this at the local level '.² GTTs may contribute to such a system; however, other sources such as incident reports, guideline/ standards compliance audits, collections of quality indicators, consumer experience feedback and coroners' reports would also be necessary for an integrated and comprehensive framework of safety and quality data in primary care.¹²

Limitations

Only five general practices were involved in this study and they were recruited by a convenience method, thereby limiting the generalisation of findings more broadly in Australia. There is likely to have been variation between medical record reviewers in the interpretation and thresholds for deciding whether an AE was present, and also in the rating of preventability and level of harm. Inter-rater reliability was not formally tested and requires more research.

Another source of potential bias is the use of reviewers examining medical records from their own practices. The option of using external reviewers was explored but clinicians expressed concern about external reviewers accessing patient-level data. For larger studies of this nature, the protection of statutory immunity may be an option.

Implications for general practice

The study found that significant levels of information about AEs exist in general practice medical records. Rates of harm are broadly in line with a similar study internationally, although with our small sample size, a national rate should not be inferred.

Table 4. Adverse events by ICPS incident type				
ICPS incident type	n	%		
Medication	29	66		
Healthcare-associated infections	6	14		
Clinical management	4	9		
Assessment/diagnosis	3	7		
Fall	1	2		
Pressure ulcer	1	2		
Total	44	100		

Table 3. Number and percentage of AEs by trigger and medical records reviewed

Trigger	Medical records in which a trigger was positive		AE				
	n ^a	%	$\mathbf{n}^{\mathbf{b}}$	%°	Positive predictive value ^d		
Three contacts in 1 week?	110	26	19	4.4	17		
Admitted to hospital?	83	19	17	4.0	20		
After-hours consult?	38	9	7	1.6	18		
Opioid analgesia?	52	12	5	1	10		
NSAID/COX ₂ inhibitors?	32	7	2	0.5	6		
eGFR ≤60 mL/min/1.73 m²?	102	24	2	0.5	2		
Benzodiazepine?	42	10	1	0.2	2		
Antipsychotic?	6	1	1	0.2	17		
Haemoglobin level ≤100 g/L?	5	1	1	0.2	20		
INR >4.5?	1	0	0	0	0		
AE, adverse event; COX ₂ , cyclooxygenase 2; eGFR, estimated glomerular filtration							

rate; NSAID, non-steroidal anti-inflammatory drug

a = number of medical records with positive triggers

b = number of medical records with an adverse event

 $c = b/428 \times 100 (428 = number of medical records reviewed)$

 $d = b/a \ge 100$

Although time taken to review records was relatively low, more research is necessary in the Australian general practice environment to determine the cost- and time-effectiveness and reliability of GTTs, and to validate their use in a larger number and variety of general practices, including those in rural and remote areas. This could lead to development and publication of an Australian general practice GTT version with associated definitions, data collection tools and training resources. The use of such a tool could be linked to the Safety and Quality (Standard 3.1) RACGP Standards for general practices (currently in 4th edition).¹⁷ GTTs are a form of clinical audit, which is an acceptable activity under the RACGP's Quality Improvement and Continuing Professional Development (QI&CPD) program.¹⁸

Tools such as GTTs should be supported by a policy and funding framework, and incentives that allow time off the 'fee-forservice treadmill', as the current cost of many high-quality safety and quality activities is potentially prohibitive. For example, in this study GPs were paid to backfill staff to undertake the reviews.

Comprehensive frameworks for systematically understanding the nature of safety and quality issues at general practice, regional and national levels, and provide incentives and resources for continuous improvement within general practice and, more broadly, within primary health care, are necessary for a deeper understanding of and response to safety priorities.

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