CLINICAL AUDIT

Improving safety for patients taking an ACE inhibitor/ARB + diuretic





Audit focus

This audit is a tool for primary care health professionals to increase medicines safety in their practice. The aim is to ensure that patients who are prescribed anti-hypertensive medicines are not put at increased risk of acute kidney injury.

Background

Angiotensin converting enzyme (ACE) inhibitors, angiotensin-II receptor blockers (ARBs), diuretics and non-steroidal anti-inflammatory drugs (NSAIDs) all have the potential to decrease renal function. When any of these medicines are prescribed together the patient's risk of acute kidney injury (AKI) is increased. ACE inhibitors/ARBs and diuretics are often taken concurrently, either as individual medicines or combination formulations. In patients taking ACE inhibitors/ARBs and diuretics particular care is required to avoid NSAIDS (the "triple whammy") due to the increased risk of AKI. The addition of a NSAID to the patient's treatment may also reduce the blood pressure-lowering effect of the ACE inhibitor and reduce the volume depleting effect of the diuretic.

When treatment with an ACE inhibitor/ARB and a diuretic is initiated, it is helpful to highlight the need to avoid NSAIDs in the patient's notes to alert other clinicians who may consider prescribing a NSAID in the future. Patients taking an ACE inhibitor/ARB and a diuretic should be warned of the risks of using NSAIDs and should be advised to avoid using over-the counter (OTC) NSAIDs, including combination products that contain NSAIDs, e.g. paracetamol and ibuprofen.

If a NSAID must be prescribed to a patient already taking an ACE inhibitor/ARB and diuretic, the lowest effective dose should be used for the shortest possible duration. Patients should also be advised to:

- Maintain adequate fluid intake at all times
- Avoid additional NSAIDs
- Manage periods of acute illness carefully, e.g. by maintaining good fluid intake, stopping the NSAID and seeking medical attention if their condition deteriorates

Before the triple whammy is initiated, a baseline measurement of serum creatinine is essential as it may be required later to diagnose AKI. A follow-up assessment, with repeat measurements of body weight, blood pressure, serum creatinine and electrolytes, within the first month of treatment may also be beneficial, due to the increased risk of AKI during this period.

Further information is available from: "Avoiding the "triple whammy" in primary care: ACE inhibitor/ARB + diuretic + NSAID", bpac.org.nz/2018/triple-whammy.aspx

Audit Plan

Summary

This audit identifies patients who are taking ACE inhibitors/ ARBs and diuretics. Patient management is then assessed to determine if appropriate steps have been taken to maximise safety, including patient advice, avoidance of NSAIDs and testing of renal function in patients who do require NSAIDs.

Recommended audit standards

Ideally, all patients who are taking an ACE inhibitor/ARB with a diuretic will have documented evidence in their patient record of a discussion about the need to avoid NSAIDs. Patients who are prescribed a NSAID at any time while also prescribed an ACE inhibitor/ARB and a diuretic, should also have a baseline renal function test recorded prior to the NSAID prescription, e.g. in the last three months, and have documented evidence that strategies to avoid the adverse effects of treatment have been discussed.

Audit Data

Eligible patients

All patients who are currently taking an ACE inhibitor/ARB and a diuretic are eligible for this audit.

Identifying patients

You will need to have a system in place that allows you to identify eligible patients. Many practices will be able to identify patients by running a "query" through their PMS. Once a patient taking an ACE inhibitor/ARB and a diuretic is identified, the patient's notes should then be reviewed to check if they have been concurrently prescribed a NSAID at any time.

Sample size

A sample size of 30 patients who are currently taking an ACE inhibitor/ARB and a diuretic is sufficient for the purposes of this audit.

Criteria for a positive outcome

A positive result is if a patient who is currently taking an ACE inhibitor/ARB and a diuretic has documented evidence in their patient record of a discussion about the need to avoid NSAID use. Ideally, the notes would also be flagged to alert other clinicians in the practice to avoid prescribing NSAIDs.

If the patient has also been concurrently prescribed a NSAID at any time they should have:

- The results of a renal function test recorded in the three months prior to the NSAID prescription
- Documented evidence that strategies to avoid adverse effects have been discussed, e.g. maintaining fluid intake, avoiding other NSAIDs and managing acute illness

Data analysis

For each patient who is currently taking an ACE inhibitor/ARB and a diuretic, record whether or not there is documented evidence in their record that the need to avoid NSAIDs has been discussed. If the patient has also been prescribed a NSAID at any time, note if a renal function test has been performed in the three months prior to the NSAID prescription and if there is documented evidence of a discussion about strategies to avoid adverse effects.

Patients who do not meet the criteria for a positive result, i.e. they have "No" recorded in any column the data sheet, should be flagged for review.

Using clinical audits for improving practice and patient outcomes

Clinical audits can be an important tool to identify where gaps exist between expected and actual performance. Once completed, they can provide ideas on how to change practice and improve patient outcomes. General practitioners are encouraged to discuss the suitability and relevance of their proposed audit with their practice or peer group prior to commencement to ensure the relevance of the audit. Outcomes of the audit should also be discussed with the practice or peer group; this may be recorded as a learning activity reflection if suitable.

The Plan, Do, Study, Act (PDSA) model is recommended by the Royal New Zealand College of General Practitioners (RNZCGP) as a framework for assessing whether a clinical audit is relevant to your practice. This model has been widely used in healthcare settings since 2000. It consists of two parts, the framework and the PDSA cycle itself, as shown in Figure 1.

1. The framework

This consists of three questions that help define the "what" and "how" of an improvement project (in this case an audit). The questions are:

- "What are we trying to accomplish?" the aim
- "How will we know that a change is an improvement?" what measures of success will be used?
- "What changes can we make that will result in improvement?" – the concept to be tested

2. The PDSA cycle

This is often referred to as the "engine" for creating, testing and carrying out the proposed changes. More than one cycle is usually required; each one is intended to be short, rapid and frequent, with the results used to inform and refine the next. This allows an ongoing process of continuous learning and improvement.

Each PDSA cycle includes four stages:

- Plan decide what the change to be tested is and how this will be done
- **D**o carry out the plan and collect the data
- Study analyse the data, assess the impact of the change and reflect on what was learned
- Act plan the next cycle or implement the changes from your plan

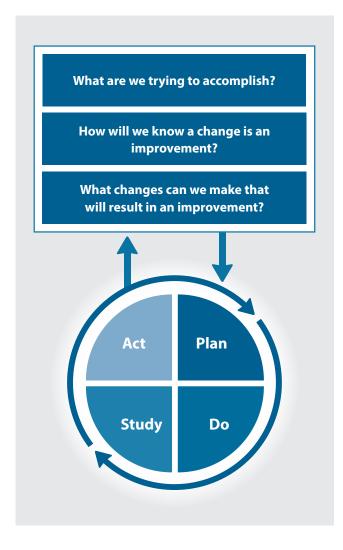


Figure 1. The PDSA model for improvement.

Source: Plan, Do, Study, Act (PDSA) cycles and the model for improvement

Claiming credits for Te Whanake CPD programme requirements

Practice or clinical audits are useful tools for improving clinical practice and credits can be claimed towards the Patient Outcomes (Improving Patient Care and Health Outcomes) learning category of the Te Whanake CPD programme, on a credit per learning hour basis. A minimum of 12 credits is required in the Patient Outcomes category over a triennium (three years).

Any data driven activity that assesses the outcomes and quality of general practice work can be used to gain credits in the Patient Outcomes learning category. Under the refreshed Te Whanake CPD programme, audits are not compulsory and the RNZCGP also no longer requires that clinical audits are approved prior to use. The college recommends the PDSA format for developing and checking the relevance of a clinical audit.

To claim credits go to the RNZCGP website www.rnzcgp.org.nz

If a clinical audit is completed as part of Te Whanake requirements, the RNZCGP continues to encourage that evidence of participation in the audit be attached to your recorded activity. Evidence can include:

- 1. A summary of the data collected
- An Audit of Medical Practice (CQI) Activity summary sheet (Appendix 1 in this audit or available on the RNZCGP website).

N.B. Audits can also be completed by other health professionals working in primary care (particularly prescribers), if relevant. Check with your accrediting authority as to documentation requirements.



PO Box 6032, Dunedin Phone 03 477 5418 contact@bpac.org.nz



Data sheet — cycle 1 Improving safety for patients taking an ACE inhibitor/ARB + diuretic

Patient concurrently	Evidence of advice to patient to avoid NSAIDs	If the patient was prescribed a NSAID at any time while also prescribed an ACE inhibitor/ARB + diuretic		A positive result, i.e.
prescribed an ACE inhibitor/ARB + diuretic		Renal function tested in three months prior to NSAID prescription Yes/No	Evidence of advice about avoiding adverse effects Yes/No	√, or flagged for review, i.e. x
	Yes/No	res/NO	res/NO	
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Data sheet — cycle 2 Improving safety for patients taking an ACE inhibitor/ARB + diuretic

Patient concurrently	Evidence of advice to patient to avoid NSAIDs Yes/No	If the patient was prescribed a NSAID at any time while also prescribed an ACE inhibitor/ARB + diuretic		A positive result, i.e.
prescribed an ACE inhibitor/ARB + diuretic		Renal function tested in three months prior to NSAID prescription Yes/No	Evidence of advice about avoiding adverse effects Yes/No	√, or flagged for review, i.e. ★
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SUMMARY SHEET

Audit of medical practice (CQI activity)

Topic:						
Improving safety for patients taking an ACE inhibitor/ARB + diuretic						
Activity designed by (name of organisation, if relevant):						
Bpac ^{nz}						
Doctor's name:						
Results discussed with peer group or colleagues?	Date:					
Yes No						
FIRST CYCLE						
DATA: Date of data collection:						
CHECK: Describe any areas targeted for improvement as a result of analysing the data collected.						
The state of the s						
ACTION: Describe how these improvements will be implemented.						
MONITOR: Describe how well the process is working. When will you undertake a second cycle?						