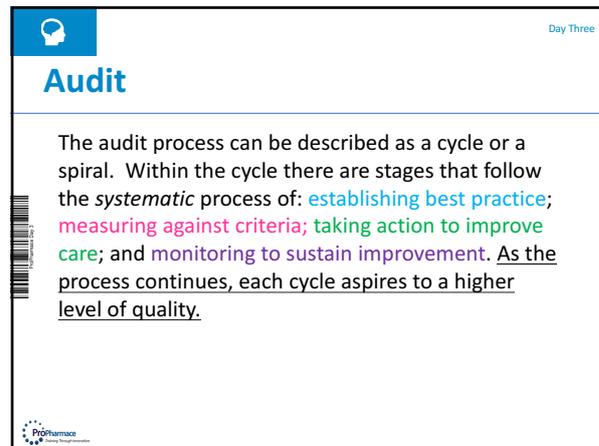




Introduction to AUDIT

ProPharmace
Pre-registration Training 2019-20

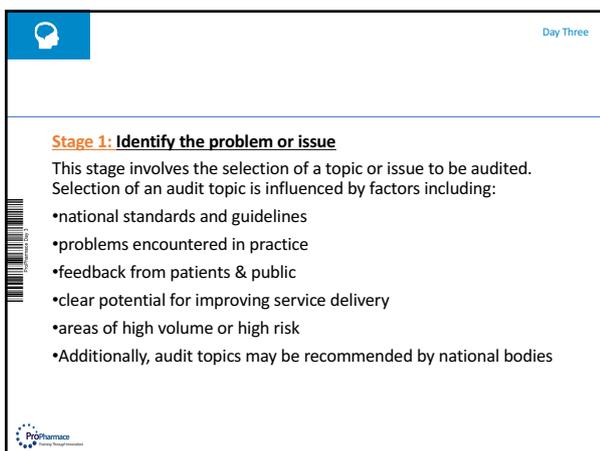
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Audit

The audit process can be described as a cycle or a spiral. Within the cycle there are stages that follow the *systematic* process of: **establishing best practice**; **measuring against criteria**; **taking action to improve care**; and **monitoring to sustain improvement**. As the process continues, each cycle aspires to a higher level of quality.

The slide includes a blue header with a white brain icon and the text 'Day Three'. A barcode is visible on the left side. The ProPharmace logo is in the bottom left corner.

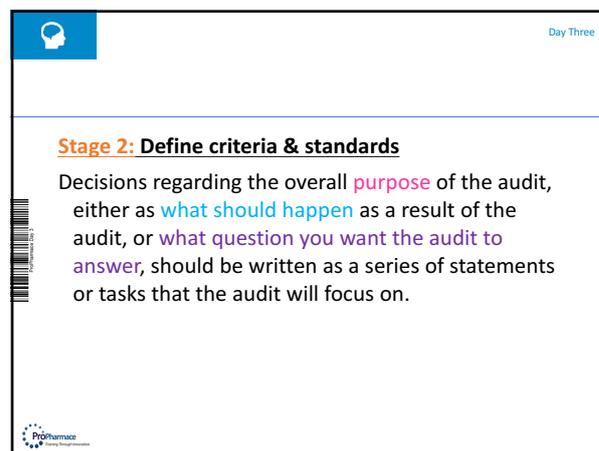


Stage 1: Identify the problem or issue

This stage involves the selection of a topic or issue to be audited. Selection of an audit topic is influenced by factors including:

- national standards and guidelines
- problems encountered in practice
- feedback from patients & public
- clear potential for improving service delivery
- areas of high volume or high risk
- Additionally, audit topics may be recommended by national bodies

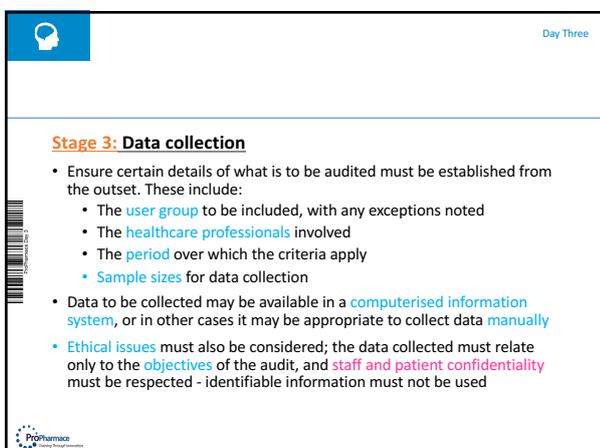
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Stage 2: Define criteria & standards

Decisions regarding the overall **purpose** of the audit, either as **what should happen** as a result of the audit, or **what question you want the audit to answer**, should be written as a series of statements or tasks that the audit will focus on.

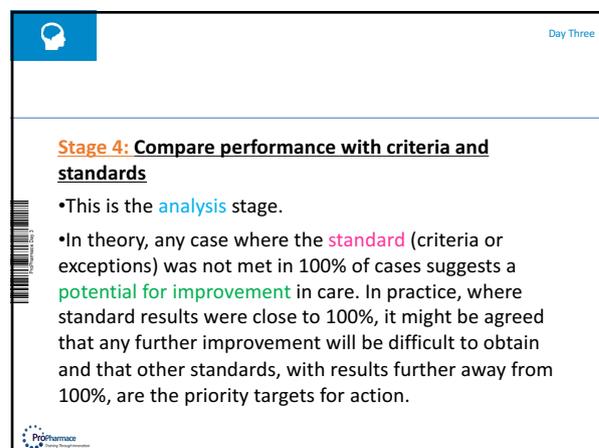
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Stage 3: Data collection

- Ensure certain details of what is to be audited must be established from the outset. These include:
 - The **user group** to be included, with any exceptions noted
 - The **healthcare professionals** involved
 - The **period** over which the criteria apply
 - **Sample sizes** for data collection
- Data to be collected may be available in a **computerised information system**, or in other cases it may be appropriate to collect data **manually**
- **Ethical issues** must also be considered; the data collected must relate only to the **objectives** of the audit, and **staff and patient confidentiality** must be respected - identifiable information must not be used

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Stage 4: Compare performance with criteria and standards

- This is the **analysis** stage.
- In theory, any case where the **standard** (criteria or exceptions) was not met in 100% of cases suggests a **potential for improvement** in care. In practice, where standard results were close to 100%, it might be agreed that any further improvement will be difficult to obtain and that other standards, with results further away from 100%, are the priority targets for action.

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Day Three

Stage 5: Implementing change

- Results of audit discussed
- Agreement about recommendations for change
- Using an **action plan** to record these recommendations is **good practice**
- Who has agreed **to do** what and by **when**
- Each point needs to be **well defined**, with an **individual named** as responsible for it, and an agreed **timescale for completion**



Day Three

Re-audit: Sustaining Improvements

- After an agreed period, the audit should be **repeated**.
- The **same strategies** for identifying the sample, methods and data analysis should be used to ensure comparability with the original audit.
- The re-audit should demonstrate that the **changes have been implemented** and that **improvements have been made**. Further changes may then be required, leading to additional re-audits.
- **Critical to achieve the standards of healthcare delivery identified in stage 2.**



Day Three

AUDIT project

- All ProPharmace Trainees are required to complete an audit project.
- Initiate a discussion with your tutor about an audit topic that would be suitable to complete at your pharmacy.





ProPharmace Audit Project Guidance

The GPhC requires all pre-registration trainees to complete an AUDIT (Performance Standard A4.8)

If you wish to submit your audit, please do so by bringing a printed copy to your Day 7 training day to hand in for marking. Your audit should satisfy the criteria detailed below.

<p><u>Headings:</u></p> <ul style="list-style-type: none">• Authors• Title• Introduction• Aims and Objectives• Participants (Consent Obtained)• Action Plan• Standards• Methods and Resources used• Results• Discussion and Conclusion• Proposals for future (Review Date)• References	<p><u>Format</u></p> <ul style="list-style-type: none">• Single line spacing and justified in both margins• Page margins: 2.0cm• Main titles - Arial font 12pt bold• Main text - Arial font 11pt• Maximum of 1000 words• All abbreviations should be explained• References: Vancouver style
<p>DEADLINE FOR SUBMISSION: Bring a paper copy of your Audit Project to your Day 7 ProPharmace Training Day</p>	

Tip: Download the PSNC Guide to Clinical Audit at <https://psnc.org.uk/contract-it/essential-service-clinical-governance/clinical-audit/>

Important:

- Many projects are inaccurately described as audits. An audit must include *standards*
- Late submissions may not be accepted; plan ahead to complete your project in a timely manner using SMART objectives
- Your audit project mark will be shared with your tutor