

# Unit 88: Preparing Lower GI Equipment: Endoanal Ultrasound

<b>Level:</b>	<b>4</b>
<b>Unit type:</b>	<b>Optional (Gastrointestinal Physiology)</b>
<b>Credit value:</b>	<b>6</b>
<b>Guided learning hours:</b>	<b>46</b>

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## Unit summary

In this unit, you will develop the understanding and skills needed to prepare equipment to investigate the lower gastrointestinal tract using endoanal ultrasound. You will also be able to undertake routine maintenance and calibration procedures on the equipment used. You will be expected to build your patient-centred professional practice and practise safely in the workplace, seeking advice and referring as appropriate to your scope of practice.

## Unit assessment requirements

There are no specific assessment requirements for this unit, however **learners completing this unit must also complete *Unit 81 Introduction to Gastrointestinal Physiology***. Please refer to the assessment strategy in *Annexe B*.

## Additional information

AC1.2 includes:

- the impact of incorrect calibration on the individual, the investigation and the results
- identification of common faults and remedial action required
- safety testing
- routine maintenance in accordance with the manufacturer's recommendations.

AC1.4 includes:

- faecal incontinence
- sepsis
- tumour identification
- sphincter dysfunction.

AC2.1 includes the:

- potential risks of using defective equipment and measures to resolve problems
- factors influencing the choice of equipment for the investigation.

AC2.2 includes:

- the requirements for the investigation environment
- ensuring that all consumables are available
- the principles of measurement for the range of equipment used
- identifying common faults and taking remedial action
- explaining the measurements that are required pre-investigation
- explaining the impact of incorrect calibration on the individual, the investigation and the results
- cleaning and disinfecting faulty catheters and documenting a manufacturer's fault report.

AC2.3 includes:

- the standard operating procedure for the equipment used
- ensuring that all consumables are available
- lubricating gel
- sheaths
- checking the patient notes for latex allergy
- checking that equipment complies with all current safety standards, e.g. BME commissioning number
- checking that equipment is safety-tested and regularly maintained in accordance with the manufacturer's recommendations
- withdrawing externally damaged transducers from individual use and completing required documentation
- using appropriate infection control and hand-washing methods
- disinfecting equipment before (and after) use according to equipment manufacturers' instructions
- applying the appropriate sheath for the purpose.

AC3.2 includes:

- disposing of any consumables safely and appropriately
- cleaning and disinfecting equipment appropriately.

## Learning outcomes and assessment criteria

To pass this unit, learners need to demonstrate that they can meet all the learning outcomes for the unit. The assessment criteria outline the requirements that the learner is expected to meet **in own area of work and in accordance with Standard Operating Procedures (SOPs)** to achieve the learning outcomes and the unit.

Learning outcomes		Assessment criteria		Evidence type	Portfolio reference	Date
1	Understand the underpinning principles of endoanal ultrasound	1.1	Explain the principles of measurement for the range of equipment used			
		1.2	Explain the purpose of equipment calibration and maintenance			
		1.3	Explain the factors influencing the choice of equipment for the investigation			
		1.4	Know the indications and contraindications for the investigation			

Learning outcomes		Assessment criteria		Evidence type	Portfolio reference	Date
2	Be able to prepare equipment and environment for endoanal ultrasound	2.1	Demonstrate the ability to select suitable equipment for the investigation in accordance with the requirements of the test			
		2.2	Prepare the environment and equipment for investigations			
		2.3	Perform the calibration of equipment at the beginning of every test according to the manufacturer's instructions			
3	Be able to clean equipment, and leave the room in a suitable condition for reuse	3.1	Demonstrate the ability to decontaminate all non-single use only equipment, and leave equipment and environment in a suitable condition for reuse, including restocking consumables			

Learner name: \_\_\_\_\_

Date: \_\_\_\_\_

Learner signature: \_\_\_\_\_

Date: \_\_\_\_\_

Assessor signature: \_\_\_\_\_

Date: \_\_\_\_\_

Internal verifier signature: \_\_\_\_\_

Date: \_\_\_\_\_

*(if sampled)*