

The Radiographic Assistant Practitioner's role in Quality Control of Radiological Equipment

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Background

Radiographic Assistant Practitioners (AP), like general support staff, are likely to be from diverse backgrounds but they differ from the general support workforce in that, as part of their duties, they perform limited clinical imaging examinations in concert with, and under the supervision of, HCPC-registered Radiographers.

The Society and College of Radiographers (SCoR) receives requests from the UK diagnostic radiography community pertaining to the scope of practice of the qualified AP. One such request is the need for guidance on the AP's role in Quality Control (QC) of radiological equipment. This briefing details such guidance and has been produced in collaboration with the Institute of Physics and Engineering in Medicine (IPEM). The Society and College of Radiographers wishes to gratefully acknowledge the support and advice of the IPEM Diagnostic Radiology Special Interest Group in the development of this guidance briefing.

Introduction

The Ionising Radiations Regulations 1999 (IRR99) 1 require that the employer makes arrangements for a suitable QC programme to be provided for all radiological equipment. The employer should consult the appointed Radiation Protection Adviser (RPA) and Medical Physics Expert (MPE) about the QC programme. There are also many guidance documents 2,3,4 containing recommended standards for the QC of equipment.

The QC programme should

- specify the frequency of testing;
- contain a written protocol detailing how to perform the testing;
- contain appropriate action level(s) for each test result;
- identify the remedial actions required in the event that the action level(s) are exceeded;
- make clear who has responsibility for carrying out the testing;
- make clear who has responsibility for acting on adverse findings.

Under IRR99, it is for the employer to decide who undertakes the testing – this can be the employer's own staff or contractors. When QC is undertaken, it is important that all other relevant aspects of IRR99 are adhered to – this includes compliance with the Local Rules and arrangements for personal monitoring.

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Under the Ionising Radiation (Medical Exposure) Regulations 2000 (IR(ME)R),⁵ anyone performing QC of radiological equipment is regarded as an IR(ME)R operator.

With their familiarity with the equipment and their knowledge of the department in which they work, APs are well placed to undertake equipment QC. Before the AP can undertake this operator duty, IR(ME)R requires the following:

- a. that the AP is entitled, in accordance with the employer's procedures, to act as an IR(ME)R operator with a scope of practice that includes the QC tests they perform and the equipment on which they perform these tests;
- b. that there is a written protocol covering the practical aspects of the QC that is performed;
- c. that the AP is adequately trained as an IR(ME)R operator, with provision made for ongoing training:
- d. that the AP, in their capacity as an IR(ME)R operator, follows the employer's procedures.

This guidance briefing outlines the steps to be taken where an employer wishes to entitle their APs to perform equipment QC.

Entitlement under IR(ME)R

Entitlement of APs to perform equipment QC should not be considered until all other steps in this guidance briefing have been completed.

The entitlement should allow the AP to be identified, whether by name or designation. It should also include their defined scope of practice, including the tests they are permitted to undertake and the equipment on which they are permitted to perform these tests. The date on which they were entitled (or dates if it varies with equipment or test) and a schedule for a review of their training records and entitlement should also be included.

Since the employer will have entitled radiographers as IR(ME)R operators, a good first step will be to amend that entitlement documentation to suit this case.

Written protocol

The protocol should include sufficient information as to allow the test set-up and technique to be accurately reproduced. It will, therefore, require detailed information on the following for each test being undertaken:

- the test equipment to be used;
- the positioning of the x-ray equipment and test equipment;
- the exposure factors;
- the result to be recorded following the test, including where to record it;
- the action level(s);
- the remedial actions required in the event that the action level(s) are exceeded;
- if the remedial action is to escalate it to senior staff, their contact details should be included;
- if the remedial action is to withdraw the equipment from use, details on how that is to be communicated to all potential users should be included;
- where to record details on the action taken in the event that action level(s) are exceeded.

The choice of action levels and remedial actions is a local decision that should be taken with expert

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advice. Usually this will have come from the RPA and / or MPE.

In compliance with IR(ME)R, all protocols require periodic review. Therefore this protocol should be subject to regular review (generally annually, though to be decided locally). It should also be reviewed when the equipment is replaced, updated or modified in any significant way or if there are concerns raised about the equipment's performance. In all cases it should be updated as necessary.

Training

Adequate training involves both theoretical and practical elements. The theoretical elements must be cross-referenced against schedule 2 of the IR(ME) Regulations to ensure adequate training as an IR(ME)R operator; much of this will have been covered in the AP's pre-qualification education as well as any further training to date. An understanding of the written protocol for QC testing of all equipment that is to be included in the AP's scope of practice is essential.

Practical elements will involve meeting pre-determined competencies. The means of demonstrating and assessing competence should be identified and documented in advance of the AP's training. Competencies must include the correct recording and interpretation of the results with respect to the action level(s) and a demonstration of what to do when action level(s) are exceeded.

When the AP has covered the theoretical and practical components he/she may then be signed off as competent. Provision must be made for on-going training to maintain competence.

Competence and ongoing training should be recorded on the AP's training matrix and signed and dated by the supervisor.

Following the employer's procedures

At this stage, an AP might be ready to be entitled by the employer or to have their scope of practice increased on an existing entitlement. As an IR(ME)R operator, they are legally responsible for the practical actions they undertake and must follow the employer's procedures at all times. The AP should be made aware of this prior to their being entitled by the employer and, as with other staff, they should know the limitations of their practice.

The SCoR expects that any AP in membership maintains their place on the College of Radiographers (CoR) Voluntary Register for Accredited APs. ^{6.7} The SCoR Accredited AP in SCoR membership is covered by the Professional Indemnity Insurance for all IR(ME)R duties providing there is evidence of adequate training, entitlement and compliance with the employer's procedures.

References

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- 7. The Society and College of Radiographers Voluntary Register of Accredited Assistant Practitioners
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