

# Introduction of Image Guided Radiotherapy into Clinical Practice

## Appendix I. SELF-ASSESSMENT QUESTIONNAIRE

This questionnaire is associated with a forthcoming IAEA publication and is designed to assist centres that plan to embark on a programme of IGRT to check that they have all the necessary requirements. By the time the first patient is to be imaged with the IGRT system the answers to all the questions should be "Yes". Where gaps are identified they will need to be corrected.

**Name of the Hospital**

**Address of the Hospital**

**Name of the main contact person and position covered**

**E-mail**

**Phone number**

**Form completed on date (day/month/year)**

**1. Does your department meet the requirements for 3-D CRT of the self-assessment questionnaire of IAEA TECDOC 1588 excluding IMRT (questions 1-49)?**

Yes      No

**2. Have one or more patient groups been identified that would benefit from IGRT?**

Yes      No

**3. Have all groups of staff had at least one year experience in the planning and delivery of 3-D CRT?**

Yes      No

**4. Has an IGRT committee including a radiation oncologist, medical physicist and radiation therapist been established to oversee the introduction of IGRT?**

Yes      No

**5. Are there sufficient radiation oncology, medical physics and radiation therapy staff to ensure that the introduction of IGRT does not compromise other radiotherapy treatment including 3-D CRT?**

Yes      No

**6. Are there satisfactory service support arrangements to ensure that the IGRT equipment can be maintained at the required level of accuracy?**

Yes      No

**7. Have all groups of staff had additional education and training in IGRT?**

Yes      No

**8. Does the department have access to a dose assessment system for IGRT dosimetry?**

Yes      No

**9. Does the department have QC expertise, methodology, and tools to maintain an IGRT service?**

Yes      No

**10. Have image acquisition protocols been developed for the anatomical sites to be treated with IGRT?**

Yes      No

**11. Have IGRT protocols been developed for the anatomical sites to be treated with IGRT?**

Yes      No

**12. Have tests been carried out to ensure that the record and verify system is capable of supporting IGRT?**

Yes      No

**13. Have commissioning tests including end-to-end tests been performed to demonstrate IGRT system capability and workflow?**

Yes      No

**14. Have contingency plans been developed in case of unavailability of IGRT?**

Yes      No

**15. Additional comments:**