

Section 1: Radiation Oncology Facility Information

1. Please enter the details for your facility below

Facility Name

Site (if applicable)

Questionnaire completed by

Contact phone number

Contact email

Date completed

Section 2: Inversely Planned Treatment Techniques

Please select those treatment techniques which will be used for patients at your facility

2. IMRT Static Gantry: step and shoot

Yes

No

3. IMRT Static Gantry: dynamic leaves

Yes

No

4. VMAT: One or more arcs

Yes

No

5. VMAT Technique

Single

Rapid

Multiple

Smart

Hybrid

Other

Other (please specify)

6. Helical Tomotherapy

Yes

No

7. State dose rate used for clinical treatments in MU/min

Section 3: Radiation Oncology Planning and Treatment Information

8. How many patients were treated at your facility with IMRT in the last year?

Head and Neck

Brain

Chest

Breast

Abdomen including Gastric

Pelvis including Prostate, Bladder and
Gynaecological

Extremities

Other (please specify type and number of patients)

9. How many patients were treated at your facility with VMAT in the last year?

Head and Neck

Brain

Chest

Breast

Abdomen including Gastric

Pelvis including Prostate, Bladder and
Gynaecological

Extremities

Other (please specify type and number of patients)

10. How many patients were treated at your facility with Helical Tomotherapy in the last year?

Head and Neck

Brain

Chest

Breast

Abdomen including Gastric

Pelvis including Prostate, Bladder and
Gynaecological

Extremities

Other (please specify type and number of patients)

11. Treatment Planning System

Manufacturer and model

Software version

Dose calculation algorithm

Typical dose grid spacing in cm

Plan data export in DICOM-RT format possible

12. Do you have an additional treatment planning system?

Yes

No

13. Treatment Planning System

Manufacturer and model

Software version

Dose calculation algorithm

Typical dose grid spacing in cm

Plan data export in DICOM-RT format possible

14. Oncology Information Management System

Manufacturer and model

Software version

15. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

Local Linac Name

Imaging tools used

EPID model eg Varian aS500, aS1000

Support arm type eg Varian R-arm or E-arm

EPID software version eg Varian IAS2, IAS3

Beam energy (MU) used for IMRT/VMAT treatments

Dose rate (MU/min) used for IMRT/VMAT treatments

EPID surface position in cm from target

EPID installation date

EPID integrated mode license

Is DIBH used for breast treatments?

Specify equipment used for DIBH

Describe imaging and verification process for DIBH

16. Do you have an additional treatment unit?

Yes

No

17. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

Local Linac Name

Imaging tools used

EPID model eg Varian aS500, aS1000

Support arm type eg Varian R-arm or E-arm

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19. Treatment Unit

Manufacturer and model

Software version

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20. Do you have an additional treatment unit?

Yes

No

21. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

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Yes

No

23. Treatment Unit

Manufacturer and model

Software version

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Serial Number

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Describe imaging and verification process for DIBH

24. Do you have an additional treatment unit?

Yes

No

25. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

Local Linac Name

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Describe imaging and verification process for DIBH

26. Do you have an additional treatment unit?

Yes

No

27. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

Local Linac Name

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Describe imaging and verification process for DIBH

28. Do you have an additional treatment unit?

Yes

No

29. Treatment Unit

Manufacturer and model

Software version

Treatment couch model

Serial Number

Local Linac Name

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Specify equipment used for DIBH

Describe imaging and verification process for DIBH

Section 4: IMRT QA Procedures

30. Describe verification of absolute dose including tolerance limits per field or total. Specify any software used for independent MU checks, with tolerances.

31. Describe verification of dose distribution, including equipment used and analysis method. For gamma analysis specify tolerance criteria eg 3%/3mm, pass rate eg 95% and threshold eg 10%.

32. Describe IGRT protocol with tolerance limits for anatomical sites including head and neck, prostate

33. Please provide a copy of local physics patient specific in-house QA protocol and form by email to qa@trog.com.au and confirm email has been sent below

- Yes, form emailed
- No, form not emailed

Section 5: External Dosimetry Audit for Facility Credentialing

TROG requires evidence of a level 1 dosimetry audit eg Australian Clinical Dosimetry Service or international equivalent.

TROG also requires evidence that your facility has participated in at least one external dosimetry audit of IMRT, VMAT and/or TomoTherapy, and will accept international audits provided the results are within TROG tolerances.

Please provide details of all external audits by recognised bodies within the last 5 years.

34. Level 1 Audit Details

Auditing body and country

Postal OR site visit

Report emailed to qa@trog.com.au (Yes/No)

35. IMRT/VMAT/TomoTherapy Audit Details

Auditing body and country

Postal OR site visit

Anatomical site

Report emailed to qa@trog.com.au (Yes/No)

36. Have you completed an additional audit in the last 5 years?

Yes

No

37. Audit Details

Auditing body and country

Postal OR site visit

Anatomical site

Report emailed to qa@trog.com.au (Yes/No)

38. Have you completed an additional audit in the last 5 years?

Yes

No

39. Audit Details

Auditing body and country

Postal OR site visit

Anatomical site

Report emailed to qa@trog.com.au (Yes/No)

Section 6: Additional Relevant Information

40. Please provide any further information which may be relevant to your participation in TROG trials

TROG values your participation in their research and is grateful for your support in completing this questionnaire. All data will be handled confidentially and only for the purpose of TROG clinical trials.

Please notify qa@trog.com.au when you have completed the survey and attach any relevant reports as requested in the questionnaire.