

# **Evaluation of Patient-Specific Quality Control (QC) for Markerless Dynamic Tumor Tracking (MDTT) Deliveries**

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### INTRODUCTION

Dynamic tumour tracking (DTT) for stereotactic ablative radiotherapy (SABR) often requires fiducial marker insertion to localize and verify target positions. The Vero4DRT (Brainlab AG) linac also offers a *markerless* dynamic tumor tracking (MDTT) module which can eliminate the need for fiducial marker insertion by tracking soft tissue targets or other anatomical surrogates. The patient-specific Quality Control (QC) results for marker-based vs *markerless* DTT deliveries can be compared.

## AIM

To report on patient-specific QC results for markerless dynamic tumour tracking (MDTT) deliveries in a phantom and compare to conventional marker-based deliveries for liver and lung SABR treatments.

### METHOD

- Vero4DRT linac (Figure 1), delivered clinical plans, 5 7 fields step-and-shoot IMRT DTT plans to two commercially available motion phantoms, modified with in-house additions.
- Motion platforms were programmed with patient-specific respiratory motion traces acquired at 4DCT.

#### Vero4DRT quick description

- DTT achieved using a gimbal-mounted waveguide and collimation system
- Motion-correlation models are built between an external IR reflector signal and internal kV fluoroscopy-detected tracking structures (implanted markers or soft-tissue landmarks)
- Verification orthogonal kV image pairs are acquired during DTT delivery (at 1 Hz frequency)
- Image-detected vs model-predicted internal positions recorded in ExacTrac imager log files

### *Ion chamber point-dose measurement* (Figure 2)

A 15 x 15 cm<sup>2</sup> acrylic phantom was placed on a Brainlab-supplied 1D moving platform. The phantom is equipped with:

1) 0.6cc farmer chamber insert

- 2) "Lung tumour" like object (imitation small bird egg) as MDTT tracking structure
- 3) 3 implanted gold-seed fiducials (1 mm x 3 mm)

### *<u>Film (Gafchromic</u> ™ <u>EBT3 2D measurement</u>* (Figure 3)

Quasar<sup>™</sup> respiratory motion phantom with an in-house acrylic cylindrical insert containing: 1) 6.0 x 7.5 cm<sup>2</sup> film receptacle (coronal plane) with 3 puncture fixtures to mark corners 2) 'Liver Dome' shaped end-cap as MDTT tracking structure

3) 3x gold-seed fiducial markers (1 mm x 3 mm)

<u>Delivery</u>: 5 clinical plans were delivered to both phantoms in the following modes: Static mode (no motion)

- Fiducial-marker DTT (conventional tracking)
- Markerless DTT (novel tracking):
  - $\rightarrow$  Tracking Structure="Lung tumour" (egg) or "Liver Dome" (acrylic cap)

### <u>Analysis</u>

- All chamber measurements were compared to RaySearch RayStation<sup>™</sup> calculated dose
- 2D film distributions from marker- and markerless- DTT were compared to static deliveries → Epson Expression 10000 XL scanner, FILMQA<sup>™</sup> PRO software, 2D gamma analysis
- Imaging log file statistics (3D vector deviations between detected vs predicted tracking structure locations) were collected for all plans.

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sert with implanted

old-seed fiducials

puncture

fixtures

gold seed



Figure 3. QUASAR motion phantom with IR respiratory pad and in-house insert. Inset: "Liver Dome" insert with film plane, puncture fixtures and fiducial markers.

### RESULTS

- markerless DTT
- See Graph 1

#### Film measurement

See Graph 2

## CONCLUSIONS

### ACKNOWLEDGEMENT



Both tracking methods meet our institutional passing criteria for patient-specific QC

marker based tracking (based on log file analysis).

• However, no impact on dose delivery is observed (2D film measurements).

Both tracking methods provide an equally viable treatment options.

dome phantom used in this study