

Introduction

MR-guided stereotactic radiotherapy (MRgRT SABR) allows for reshaping the radiation dose based on daily changes in the shape, size and position of the tumour and surrounding healthy anatomy. This presents a radical change in the radiotherapy workflow and the additional steps can result in treatment times of over an hour for many cases.

Decreasing treatment times is a key driver in the growth of MRgRT SABR. To achieve this and quantify changes, detailed measurement of the time spent on each step is required. Moreover, measuring the effect of new technologies and processes is important in understanding their success.

Timing data

The treatment steps involved in MRgRT are listed in Table 1. Analysis of long file data from the treatment console allows for precise measurement of each of these stages.

Data for over 1000 treatment fractions delivered from December 2019 to June 2021 was available for review. This volume of data is able to give a comprehensive review of the time taken to deliver treatments during this period. Focus was given to the three major sites treated: prostate, pancreas and liver.

Workflow element	Steps involved				
Setup imaging	Low- and high-resolution imaging Image review and matching				
Target contouring	Electron density overrides Target review and recontouring				
OAR contouring	OAR review and recontouring				
Independent QA 1	Tracking structure delineation Physics review				
Re-optimisation	Dose re-optimisation, review and sign-off				
Independent QA 2	Physics review of dose Independent dose calculation				
Verification imaging	Verification of target position Adjustment of tracking parameters				
Treatment delivery	Beam on and MLC movement Adjustment of tracking parameters				

Table 1: The precise time of user interactions with the treatment machine used to mark specific point in the treatment process. From this data, the elapsed time for each step can be extracted.

A time and motion study of stereotactic MR-guided adaptive radiotherapy (SMART) on a 0.35 T MR-Linac

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Comparison of treatment sites

The overall median treatment time for prostate cases is shorter than for pancreas and liver cases. However, not all workflow steps contribute to this difference. Organ at risk (OAR) contouring and beam on time are longer in pancreas and liver treatments. Whereas beam on time is shorter in prostate cases.

Step imaging: In pancreas and liver cases, 17-25 second breath-hold imaging is used. Whereas higher resolution images are acquired for prostate cases. This noticeably impacts the time of the setup imaging step.



Contouring: The target contouring step is of similar **Beam on time:** Due to the gated treatment delivery, beam on time is longer in pancreas and liver cases. In extreme length for all treatment sites. However, OAR contouring is significantly longer for more complex pancreas and cases where patients struggle to repeatedly hold their breath, beam on time can approach 40 minutes. liver cases.

A key factor in efficient scheduling of treatments is to reduce the case-to case variation in treatment time. Common variation of over fifteen minu per fraction is seen for pancreas patients (Table 2).

Greatest variation for individual workflow steps is seen in OAR contouri (Table 3) and beam on time. Ensuring regular review and feedback contouring ensures consistency and reduces variations in workflow timings. High-speed MLCs, patient coaching and improved tracking and gating can reduce variation in this step.

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utes	Treatment site	25th percentile	Median	75th percentile		
	Prostate	42.8 minutes	48.0 minutes	54.3 minutes		
ring on	Pancreas	52.1 minutes	60.0 minutes	69.2 minutes		
	Liver	50.4 minutes	58.5 minutes	64.8 minutes		

Table 2: 25th percentile, median and 75th percentile times for overall treatment for each site.

For prostate cases, treatment times have remained consistent since the service started in December 2019. Pancreas and liver treatments show greatest variation in individual fraction timings. For liver cases, changing the way in which OARs are contoured to remove the need to define individual parts of GI-tract has led to more consistent treatment times.



Contouring time for individual clinicians

The time taken for OAR contouring can be significantly impacted by the attending clinician, with a variation of over 50% observed. This is possibly due to differences in site specialities, experience and confidence.

To decrease the variation seen in contouring time between different clinicians, further training focused on non-specialist areas is planned.

Workflow improvements over time

atment site	Dr A	Dr B	Dr C	Dr D	Dr E	Dr F	Dr G	Dr G
state contouring time (minutes)	9.08	7.68	6.81	6.78	10.73	6.99	8.82	8.28
other sites contouring time (minutes)	16.59	14.07	12.42	11.06	19.45	12.74	13.78	16.00

Table 3: Median OAR contouring times for eight different clinicians, separated into prostate and all other sites. Fastest and slowest times in each category are highlighted.