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Tackling the Real Problem: The Dosimetric Unknowns in Patient Setup

Treatment planning is performed on a static CT dataset, capturing a single snapshot. The resulting plan is then delivered to the patient over multiple weeks, relying on efforts to recreate that initial anatomy and setup for every fraction.

Surface Guided Radiation Therapy (SGRT) technologies have become invaluable in helping position patients consistently, using surface metrics to guide alignment. Yet, typical SGRT tolerances often set generically at 3–5 mm and 3 degrees — are rarely customised to individual patients or specific anatomical sites. These thresholds do not reflect the actual dosimetric consequences of deviations for a given patient. Patients with limited mobility introduce additional challenges in reproducibility. Even with careful setup, small positional inconsistencies can result in subtle, unintended dose shifts. In some cases, evidence of excessive or misplaced dose to normal tissues is only discovered after treatment has been delivered, when the opportunity to intervene has passed.

Patients, after all, are not phantoms, and Patient Specific Quality Assurance (PSQA) activities lose clinical meaning when a surrogate phantom stands in for the patient for independent secondary calculations and the IMRT QA process. After all the checks and approvals are done and a patient is prepared for the start of treatment, their anatomy has already changed, and it continues to change throughout the course of treatment. Breast tissue can swell or develop seromas. Abdominal organs fluctuate with varying gas or solid content. The pelvis undergoes daily shifts with bladder or rectal fill-

ing. Even with Image Guided Radiation Therapy (IGRT) to align to internal anatomy and minimise gross setup errors, insight into how day-to-day anatomical changes impact the actual dose delivered is still lacking. So the question remains:

- How do we adapt and personalise our SGRT and IGRT setup tolerances based on the true dosimetric situation for each patient?
- How does moving to more individualised tolerances help clinicians decide when it is safe to treat versus when to adapt or replan?

The Challenge Is Recognised — But Not Universally Solved

Unseen dosimetric uncertainty is now acknowledged across the field. Sophisticated platforms like the Elekta Unity MR-linac offer online adaptive radiotherapy workflows such as Adapt to Position (ATP), which aligns the day's image to the reference via rigid registration and updates the plan accordingly. However, these solutions require significant infrastructure. Many clinics lack such technology, yet still face the same challenges of anatomy-driven dose variability.

Where RadCalc Is Heading

At LAP, we are addressing this gap by building on our core strengths: precise alignment and independent dose verification. LAP lasers and LUNA 3D offer continuous, non-invasive surface tracking from simulation through delivery. Paired with

The vision: Integrating SGRT and Dose Evaluation Workflows

Pre-treatment dose evaluation

Intra-treatment QA Post-fraction analysis

Using RadCalc's fast Monte Carlo engine on the day's anatomy — asynthetic CT constructed from the original planning CT, updated with position and surface from LUNA 3D and internal details from radiographic imaging — clinics can assess the dosimetric impact before beam-on.

Rapid DVH metrics and dose difference checks provide consoleside feedback to therapists, while physicists can review remotely. This enables quick decisions on whether to proceed, adjust, or reschedule treatment before unintended doses are delivered.

By integrating LUNA 3D's continuous surface tracking with RadCalc's time-resolved, sub-second, pre-computed particle-based fast Monte Carlo dose engine, we envision accumulating dose live during treatment.

If motion occurs within the SGRT tolerances but outside the bounds of clinical acceptability — for example, in unmonitored regions or in ways that impact internal geometry — the system could trigger a dosimetric alert or even pause the beam, delivering an unprecedented intrafraction safeguard.

Combining 3D EPID in-vivo measurements on the patient's anatomy of the day with accumulated dose summation builds a comprehensive picture of the true delivered dose across all fractions.

This supports offline adaptive decisions and enhances BED-based tools to achieve accurate dose summation across complex, multicourse or multi-site treatments, ensuring gradual anatomical changes are properly accounted for throughout the treatment process.

Figure 1. Integration of the SGRT and Dose Evaluation Workflows.

RadCalc's 3D QA suite, we are creating an integrated workflow to make dosimetric insight patient-specific and actionable. RadCalc's EPID dosimetry already allows clinics to assess the dose delivered retrospectively. However, proactive insight is needed before or during treatment, not just afterwards.

Redefining QA: Integrated SGRT and Dose Evaluation Workflows

RadCalc's future centres on combining SGRT and adaptive QA through three key workflows, as shown in the vision presented in Figure 1.

A Smarter, Patient-Centric QA Standard

Together, these tools help clinics move from fixed tolerances to a personalised, dosimetrically grounded approach — no MR-linac required. With RadCalc, QA becomes dynamic, data-driven, and built around the patient, not the plan.

Let's connect and explore how this evolution in QA can support your clinic's vision!

Note: German Patent Application No. 10 2023 115 102.9, PCT Patent Application No. EP2024/065566, US Patent Application No. 19/144,366



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