





Dosimetric Impact of Anatomical Change on VHEE and VMAT Plans: A Comparative Robustness Study

VHEE'25

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Prior Work & The Knowledge Gap



Hypothesis: VHEE is more robust to anatomical change

Why that's plausible?

Simulations: MC (DesRosiers, Papiez, others) → VHEE less sensitive inhomogeneities.

Experimental: Lagzda et al. → VHEE profiles **stable** across inserts vs photons/protons.

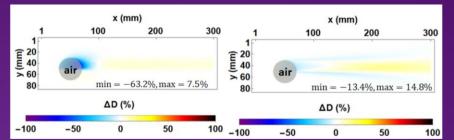
Treatment Planning: High-quality VHEE plans; **can outperform VMAT** in plan quality & OAR sparing (on static anatomy).

Gap: No direct VHEE vs VMAT under real patient changes.

This work: Proof of concept - Patient-based robustness comparison under clinically observed change.

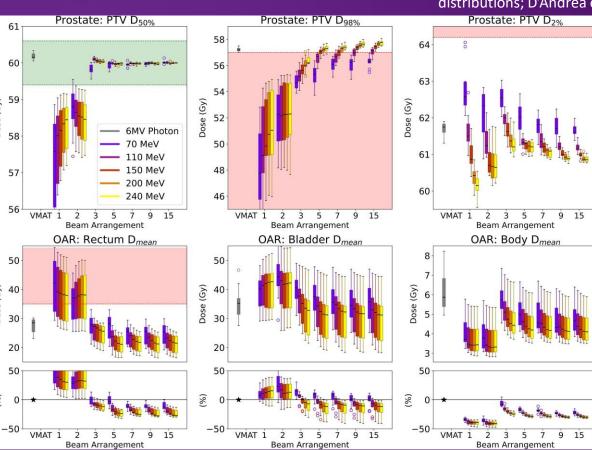
[1] Lagzda et al. Influence of heterogeneous media on Very High Energy Electron (VHEE) dose penetration and a Monte Carlo-based comparison with existing radiotherapy modalities. Nuclear Instruments and Methods in Physics Research B (2020). https://doi.org/10.1016/j.nimb.2020.09.008

[2] D'Andrea et al. Comparative treatment planning of very high-energy electrons and photon volumetric modulated arc therapy: Optimising energy and beam parameters. Physics and Imaging in Radiation Oncology (2025).



Left: Photons vs VHEE - VHEE dose profiles remain comparatively stable across air inserts (Lagzda et al., 2020; adapted [1]).

Bottom: Prostate parameter study (n=10) - boxplots across beam numbers and energies comparing 6 MV VMAT with VHEE (420 dose distributions; D'Andrea et al., 2025 [2]).





Why does this matter?



Plans built on a planning CT; delivered over many weeks:

Anatomy changes (inter- & intra-fraction):

- Tumour regression (shrinkage, oedema, baseline drift)
- OAR volume change & motion (bladder filling, rectal gas, bowel peristalsis)
- Patient weight / body-contour change

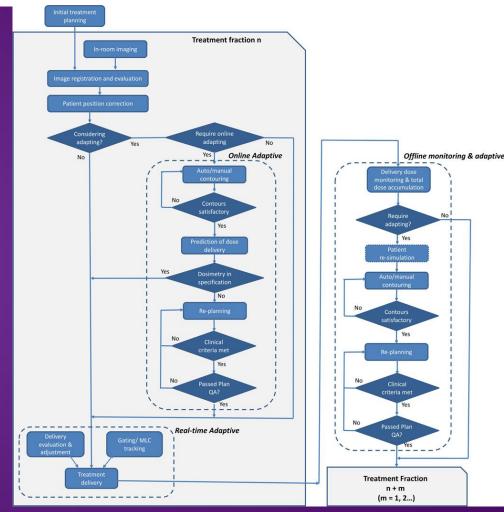
Quick implications:

- Risk: CTV underdose : ↓ local control
- **Risk:** OAR overdose: ↑ toxicity / limits breached

Action: IGRT/ART triggers → repeat imaging, plan adapt, re-fit devices

- Resource intensive
- May introduce treatment break / continue on inferior plan while replanned.

A modality with intrinsic robustness → less reliance on ART



Adaptive radiotherapy workflow showing online, offline, and real-time adaptation across fractions (imaging, registration, contouring, dose prediction/accumulation, re-planning, and plan QA). Adapted from Glide-Hurst (2021).



Method: Cases & Anatomical Change



Retrospective, preliminary, proof-of-concept cohort (n=4)

→ each triggered mid-course replan

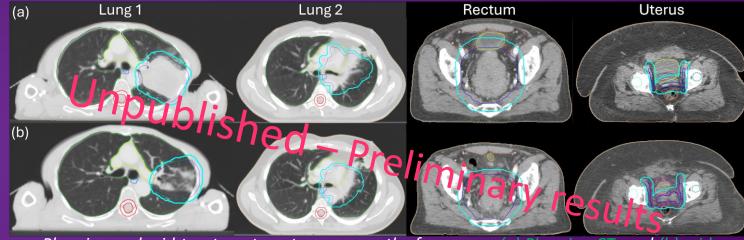
Clinical VMAT context: Approved plans; dosimetric review; replanned

Thorax

- Lung1 (60 Gy): marked regression; re-aeration.
- Lung2 (55 Gy): regression.

Pelvis

- Uterus (46 Gy): CTV stable; Bladder -67%, rectum -46%
- Rectum (45 Gy): PTV -24%, bladder -32%; Bowel stable



Planning and mid-treatment anatomy across the four cases. (a) Planning CTs and (b) midtreatment CTs for Lung1, Lung2, Rectum, and Uterus. The PTV is shown in light blue; other contours denote the re-contoured regions of interest used for evaluation



Method: Treatment Planning

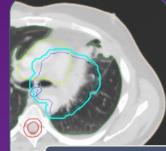




Plans

- VMAT Clinical 6MV
- VHEE: PBS 200 MeV

Planning CT



Mid-treatment CT

Recalc (noadapt)

 Fixed-MU & spotweights on mid-CT

- Target & Key OARs
- Pass/Fail vs constraints
- Δ = mid plan

Metrics & Δ

Compare

- VMAT should degrade
- Does VHEE degrade less?
- Mode of failure

	VMAT	VHEE
Delivery	Clinical 6 MV arcs (as-delivered, approved)	200 MeV, PBS, static fields
Geometry	Pelvis: full arcs Thorax: half-arc (ipsilateral)	Thorax: $n = 5$ beams, equidistant over 200° Pelvis: $n = 7$ beams, equidistant around 360°
${\bf Spot~size}$	_	$\sigma = 4 \mathrm{mm}$; spacing 1.5σ ;
Validation	Clinical TPS; dosimetrically reviewed & approved	MC (GATE/Geant4); validated (prev. work) Single-spot $\gamma \geq 98\%$ (1%/1 mm)

Redacted



Results: Pelvis exemplar - Uterus



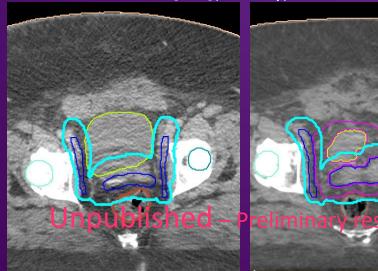
Scenario: Weight-loss, bladder –67%, rectum –46%

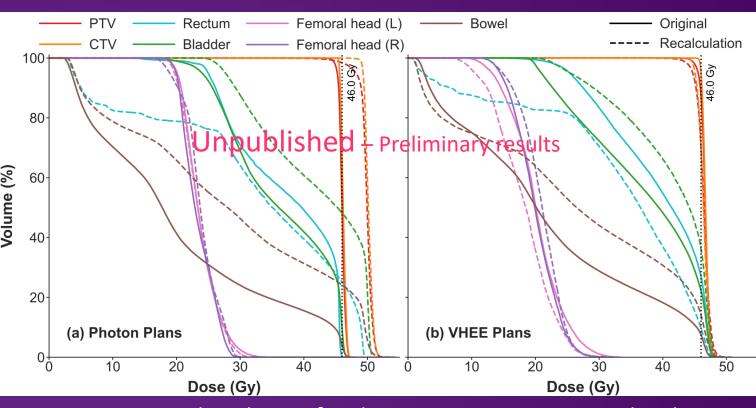
VMAT: (Plan \rightarrow Mid, tolerance):

- PTV D95 [% Rx] (\geq 97%): 98.4 \rightarrow 104.7 (pass)
- PTV D1cc [%] (≤ 110%): 102.5 → 113.3 (fail)
- Bowel V100 [cm³] (advisory): 51.3 \rightarrow 259 .3↑↑

VHEE: (Plan \rightarrow Mid, tolerance):

- PTV D95 [% Rx] (\geq 97%): 99.0 \rightarrow 97.2 (pass)
- PTV D1cc [%] (≤ 110%): 104.6 → 107.4 (pass)
- Bowel V100 [cm³](advisory): 96.5 \rightarrow 150.1 \uparrow





Uterus: combined DVHs for planning-CT optimisation and midtreatment recalculation.



Results: Thorax exemplar – Lung 1



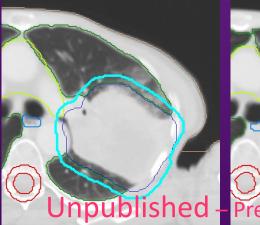
Scenario: ≈34% tumour regression + lung re-aeration;

VMAT (Plan \rightarrow Mid, tolerance):

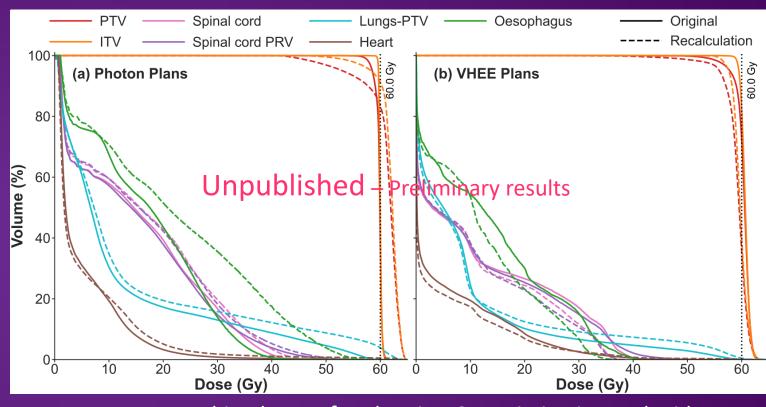
- PTV D95 [% Rx] (\geq 90%): 97.5 \rightarrow 88.4 (fail)
- PTV D1cc [%] (≤ 107%): 102.20 → 107.7 (fail)
- Cord PRV Dmax¹cc [Gy] (≤ 48): 44.5 \rightarrow 49.4 (fail)
- Oesophagus Dmean [Gy] (advisory): 16.9 → 22.2
- Lung V20 [%] (≤ 35): 17.3 \rightarrow 19.5 (pass)

VHEE (Plan \rightarrow Mid, tolerance):

- PTV D95 [% Rx] (\geq 90%): 96.3 \rightarrow 93.1 (pass)
- PTV D1cc [%] (\leq 107%): 105.4 \rightarrow 105.2 (pass)
- Cord PRV Dmax¹cc [Gy] (≤ 48): 44.4 $\rightarrow 36.2$ (pass)
- Oesophagus Dmean [Gy] (advisory): 13.3 → 11.6
- Lung V20 [%] (\leq 35): 10.2 \rightarrow 12.5 (pass)







Uterus: combined DVHs for planning-CT optimisation and midtreatment recalculation.



Clinical takeaways & next steps



Overall: In this small stress test, results suggest VHEE was more robust than VMAT.

What could this mean?

- Fewer unscheduled replans and clinical interventions
- Potential for tighter PTV margins (site-selective)
- Higher delivery confidence? (FLASH etc.)

Limitations:

- Retrospective, small cohort (n=4); proof-of-concept.
- Comparator VMAT plans were standard, not robustoptimised

Next steps:

- Larger cohorts across more sites
- Benchmarks vs robust-optimised VMAT and protons (both transmission & SOBP?)

Case	VMAT outcome	VHEE outcome
Pelvis — Uterus	! Breach: PTV $D_{1cc} = 113.3\%$ ($\leq 110\%$); $V_{100}^{\mathrm{bowel}}: 51 \rightarrow 259 \mathrm{~cm}^3$	✓ Met: $D_{95}: 99.0 \rightarrow 97.2\%$ (≥ 97%); $D_{1cc} = 107.4\%$; $V_{100}^{\text{bowel}}: 97 \rightarrow 150 \text{ cm}^3$
Pelvis — Rectum	✓ Met: coverage OK; large OAR drift $(V_{100}^{\text{bowel}}:$ $30.6 \rightarrow 280, \ V_{100}^{\text{bladder}}:$ $6 \rightarrow 42 \text{ cm}^3)$	$126, V_{100}^{\mathrm{bladder}} : 29 \rightarrow 21 \text{ cm}^3$
Thorax — Lung 1	blished – Preliming Preach: PTV $D_{95}: 97.5 \rightarrow 88.4\%$ $(\geq 90\%);$ $\operatorname{cord} PRV D_{1cc}^{\max}: 44.5 \rightarrow$ $49.4 \text{ Gy } (\leq 48); \text{ hotspot}$ $= 107.7\% \ (\leq 107\%)$	$D_{95}: 96.3 \rightarrow 93.1\%$ ($\geq 90\%$); cord PRV $D_{1cc}^{max}: 44.4 \rightarrow$
Thorax — Lung 2	! Breach: $D_{95}: 98.9 \rightarrow 92.3\%$ ($\geq 95\%$); hotspot = 110.2% ($\leq 107\%$)	! Minor: hotspot = 110.0% ($\leq 107\%$); coverage preserved $99.4 \rightarrow 99.6\%$







Thanks for listening!

Questions?

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References:

[1] Lagzda et al. Influence of heterogeneous media on Very High Energy Electron (VHEE) dose penetration and a Monte Carlo-based comparison with existing radiotherapy modalities. Nuclear Instruments and Methods in Physics Research B (2020). https://doi.org/10.1016/j.nimb.2020.09.008

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[3] Glide-Hurst et al. Adaptive Radiation Therapy (ART) Strategies and Technical Considerations: A State of the ART Review From NRG Oncology. International Journal of Radiation Oncology • Biology • Physics (2021). https://doi.org/10.1016/j.ijrobp.2020.10.021