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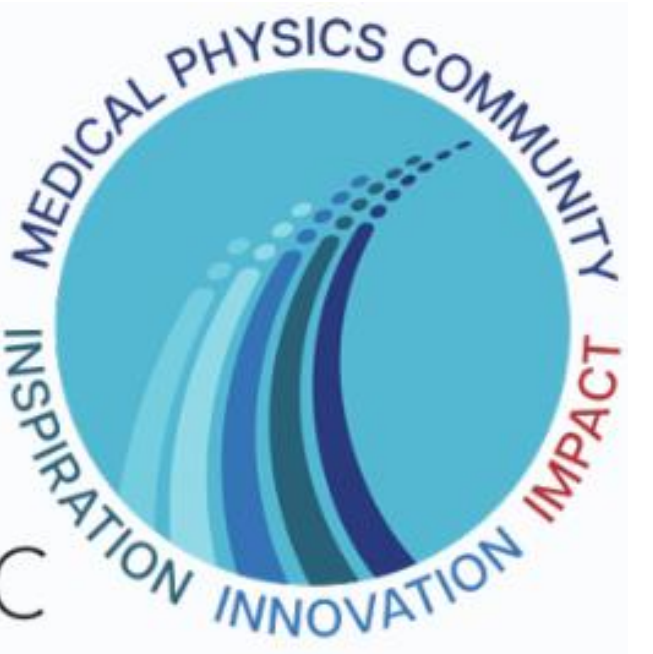
Optimizing Proton FLASH in Tangential Breast Chest Wall Radiotherapy

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Background and Aims

Post-mastectomy chest-wall (CW) irradiation is well suited for proton FLASH due to the predominance of normal tissue within the target and the feasibility of single-energy tangential transmission beam (TB) delivery. This study investigates optimization strategies to enhance FLASH activation and modeled biological sparing for large CW targets. .

Results

Clinically acceptable chest-wall coverage was achieved using a single-energy tangential TB paired with a supplemental en face clinical beam, while maintaining a simple grid-based spot layout compatible with FLASH delivery (Figure. 1a–c). Spatially, TB delivery tends to produce non-uniform FLASH activation across the chest wall, mainly towards the posterior lung region. In contrast, changing parameters such as scanning speed (V_y) and split-field TB (STB) improve the uniformity of FLASH-activated areas across both the CW-CTV and skin (Figure. 1d–k). Across the cohort, the dominant drivers of FLASH coverage were higher vertical scan speed (V_y) and STB delivery, which increased the percentage of voxels meeting FLASH criteria for both CW-CTV and skin. The highest coverage consistently occurred with the STB-ADR configuration (Figure. 2a–b). FLASH activation was model-dependent, with the ADR model achieving higher FLASH than FEM across all delivery configurations and scan speeds (Figure. 2a–b). The magnitude of sparing, evaluated by examining reductions in Δ CTV Dmean, was most pronounced in STB with ADR. GA-based sequencing produced comparable Δ CTV Dmean while improving delivery efficiency (Figure. 2c). The most promising and best outcomes of this study in terms of FLASH metrics, including maximum CW-CTV and skin FLASH activation, the most significant Δ CTV Dmean reduction, and scan-time reduction, are summarized in Table.

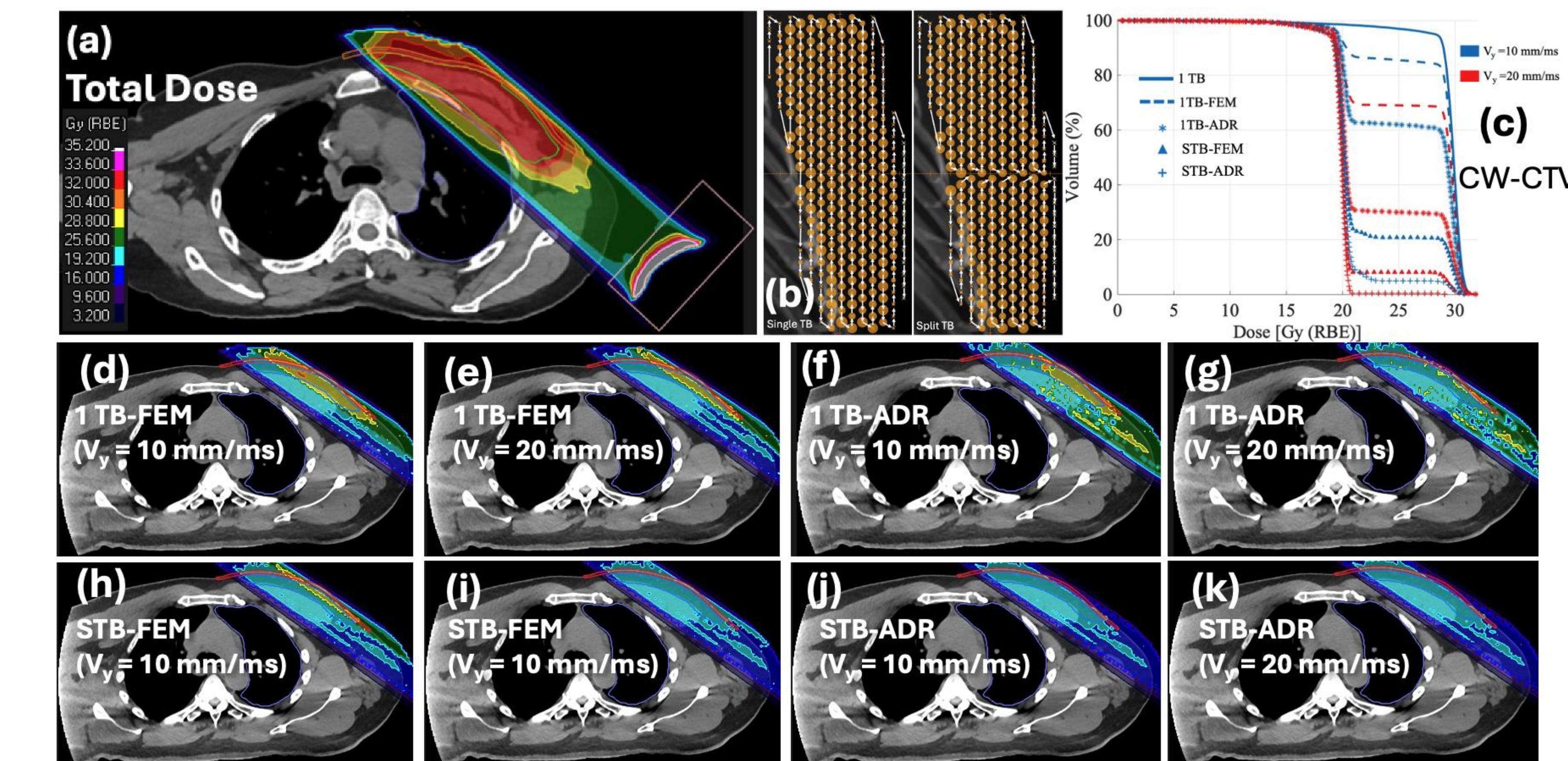


Figure 1

Conclusion

FLASH activation increased with higher scan speed ($20 > 10$ mm/ms), split-beam delivery (STB > single-TB), and ADR vs FEM, reaching 94.84% in CW-CTV and 71.96% in skin (STB, ADR at 20 mm/ms). Modeled FLASH biological sparing reduced the effective CW-CTV Dmean to normal tissue by 2.27 to 9.2 Gy, without compromising coverage. GA-based sequencing reduced scan time by 6.1%. Averaged across five patients, effective CW-CTV Dmean decreased from 29.20 Gy (TB) to 21.55 Gy (ADR-GA), and skin Dmean decreased from 20.94 to 17.42 Gy. Split-TB and higher scan speed produced the greatest FLASH activation. GA-based spot sequencing achieved FLASH coverage comparable to the split-beam delivery. A larger cohort study is needed for further validation.

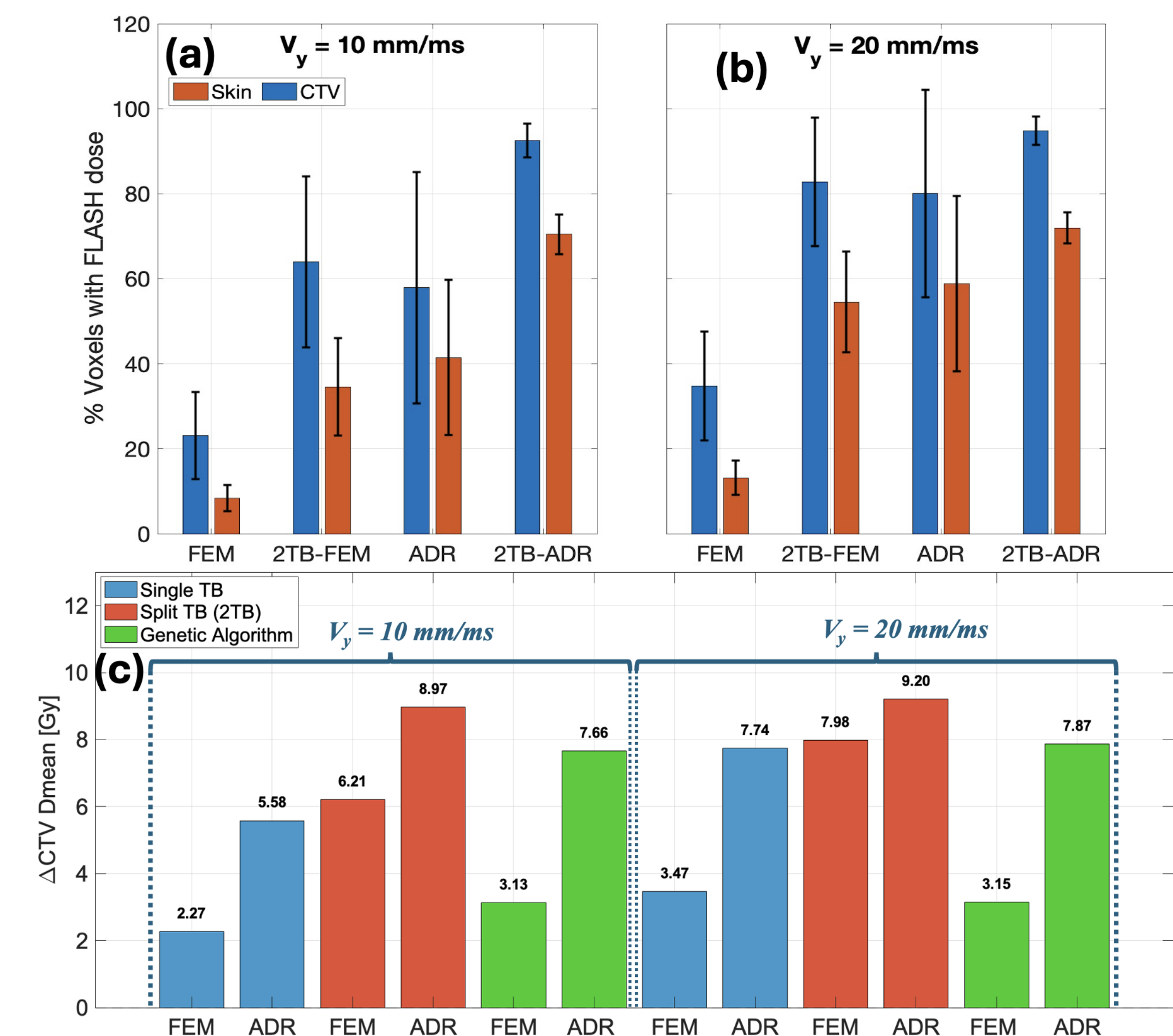


Figure 2

Metric	Best Result	Condition
FLASH-activated voxels in CTV	Up to 94.84%	STB with ADR, $V_y=20$ mm/ms
FLASH-activated voxels in Skin	Up to 71.96%	STB with ADR, $V_y=20$ mm/ms
CTV Dmean reduction	Up to ~9.2 Gy (avg), 9.57 Gy (case max)	Split TB, ADR ($V_y=20$ mm/ms)
Scan time reduction	About 6.1%	GA-sequencing vs $V_y=10$ mm/ms