

# Optimizing Treatment Efficiency In Prostate VMAT: A Dosimetric Comparison Of 10MV Flattening Filter (FF) Vs. Flattening Filter-Free (FFF) Beams

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## Abstract

*This prospective multi-institutional study evaluated 10 MV Flattening Filter-Free (FFF) versus conventional Flattening Filter (FF) VMAT for localized prostate cancer (T1c-T2c, n=13) using paired plans with identical RTOG-0815 constraints (72 Gy/30 fx) and Monte Carlo dose calculation. Comprehensive quality assurance confirmed gamma pass rates >95% (3%/2mm).*

*Results demonstrated FFF's superiority: significantly improved homogeneity (HI:  $0.08 \pm 0.02$  vs.  $0.11 \pm 0.03$ ;  $p=0.005$ ) and conformity (CI:  $0.97 \pm 0.01$  vs.  $0.95 \pm 0.02$ ;  $p=0.01$ ), reduced rectal V65Gy by 10.3% ( $21.8 \pm 5.2\%$  vs.  $24.3 \pm 6.1\%$ ;  $p=0.003$ ), bladder V65Gy by 17.1% ( $8.7 \pm 4.9\%$  vs.  $10.5 \pm 5.3\%$ ;  $p=0.02$ ), and femoral mean doses ( $1538 \pm 186$  vs.  $1637 \pm 225$  cGy;  $p=0.04$ ), alongside 26% faster delivery ( $1.7 \pm 0.2$  vs.  $2.3 \pm 0.3$  min;  $p<0.001$ ) with 31% fewer monitor units ( $880 \pm 30$  vs.  $1280 \pm 50$ ;  $p<0.001$ ).*

*These findings establish 10 MV FFF VMAT as a clinically superior, resource-efficient standard for prostate radiotherapy, simultaneously optimizing dosimetric precision, patient safety, and workflow capacity.*

**Keywords:** Prostate cancer, Flattening Filter-Free (FFF), VMAT, Dosimetric quality, Treatment efficiency, OAR sparing, Toxicity reduction

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## I. Introduction

Volumetric Modulated Arc Therapy (VMAT) has become the gold standard for prostate cancer radiotherapy, with demonstrated superiority in target coverage and organ-at-risk sparing [1-3]. Our previous study [4] established that prostate VMAT achieves excellent PTV coverage (D95 >98%) while maintaining rectal V70 <15%. However, treatment efficiency remains a significant challenge in high-throughput clinical environments.

The introduction of Flattening Filter-Free (FFF) beam technology has revolutionized linear accelerator performance, offering:

- Enhanced Dose Rates:
  - Conventional 10MV-FF: 600 MU/min
  - 10MV-FFF: 2400 MU/min (4× increase) [5,6]

- Potential Clinical Benefits:

- 40-60% reduction in beam-on time [7-9]
- Decreased intrafraction motion errors [10]
- Reduced machine component wear [11]

Despite these advantages, critical questions remain unanswered for prostate VMAT:

- Dosimetric Impact:
  - PTV homogeneity at ultra-high dose rates [12]
  - Dose gradient trade-offs for OARs [13]

- Clinical Implementation:
  - Real-world efficiency gains [14]

- Compatibility with multi-arc protocols [15]

This prospective study provides the first comprehensive evaluation of 10MV-FF versus 10MV-FFF for prostate VMAT by analyzing:

- Identical patient cohort (n=13 intermediate-risk) from [4]
- Standardized planning platform (Eclipse v16.1, AAA algorithm)
- Novel efficiency metrics:
  - Beam-on time savings
  - MU efficiency (MU/Gy)
  - Estimated patient throughput increase

The findings will establish evidence-based guidelines for FFF adoption in prostate radiotherapy, particularly relevant for:

- High-volume centers
- Resource-limited settings
- Hypofractionation protocols

## 2. Study Objectives

**1. Primary Objective:** To quantitatively compare the dosimetric performance of 10MV-FF versus 10MV-FFF beams in prostate VMAT by assessing:

- PTV coverage (D95, D98, Dmax)
- OAR doses (rectal V50, bladder V65, femoral head  $D_{max}$  and  $D_{mean}$ ) (Supported by: [4,12,13])

**2. Secondary Objectives:**

- **Efficiency Analysis:**
  - Measure beam-on time reduction (%) with FFF [7-9]
  - Calculate MU efficiency (MU/Gy) [5,6]
- **Clinical Impact:**
  - Estimate potential patient throughput increase [14]
  - Evaluate workflow integration challenges [15]

**3. Exploratory Objective:** Identify optimal cases for FFF adoption based on:

- Tumor volume (prostate ± seminal vesicles)
- Patient anatomy (rectal/bladder filling) (Preliminary data from [10,11])

## II. Materials And Methods

**1. Patient Selection** 13 intermediate-risk prostate cancer patients

- Staging: T2b-T2c
- Pathology: Gleason score 7 (3+4)
- PSA: 10-20 ng/ml
- Prostate Volume: 75-225 cc
- Prescription: 72 Gy in 30 fractions (2.4 Gy/fraction)

**2. Simulation & Immobilization** CT scanner

- Scan Range: L2 vertebra → mid-thigh
- Slice Thickness: 2.5 mm
- Position: Head-first supine
- Immobilization: knee/ankle supports
- Bladder Protocol: 300 ml water 30 mins
- Rectum Protocol: Empty

**3. Target Delineation**

- CTV: Prostate + proximal 1 cm seminal vesicles (no margin)
- PTV: CTV + 0.5 cm margin (0.3 cm posteriorly)
- OARs: Rectum, bladder, femoral heads contoured per RTOG-0815

**4. Treatment Planning**

- Linear Accelerator: Varian TrueBeam
- Treatment Planning System: Eclipse v16.1 (AAA algorithm)
- MLC: High-Definition 120 (2.5 mm leaves)

- Beam Configurations:
  - 10MV-FF: 600 MU/min
  - 10MV-FFF: 2400 MU/min
- Arc Geometry:
  - Arc 1: 179°→181° (CCW, collimator 30°)
  - Arc 2: 181°→179° (CW, collimator 330°)
- Dose Constraints:
  - PTV: V72Gy  $\geq$  95%
  - Rectum: V50Gy  $<$  25%
  - Bladder: V65Gy  $<$  15%
  - Femoral Heads max  $<$  50 Gy
- Plan Quality Homogeneity Index (HI):  $<0.1$  Conformity Index (CI):  $> 0.95$

### 5. Statistical Analysis

- Paired t-tests ( $p < 0.05$ )
- Bonferroni correction
- IBM SPSS v28

## III. Results And Discussions

**1. Patient Characteristics** This paired dosimetric analysis included 13 prostate cancer patients, with each receiving two VMAT plans:

- 10 MV Flattening Filter (FF) plan
- 10 MV Flattening Filter-Free (FFF) plan

### 2. Dosimetric Comparison

#### A. PTV Coverage: Table 1. Comparative Dosimetric Results

| Parameter         | 10 MV FFF       | 10 MV FF        | p-value  |
|-------------------|-----------------|-----------------|----------|
| D95% (%)          | 99.8 $\pm$ 0.3  | 99.1 $\pm$ 0.9  | $<0.001$ |
| Homogeneity Index | 0.08 $\pm$ 0.02 | 0.11 $\pm$ 0.03 | 0.005    |
| Conformity Index  | 0.97 $\pm$ 0.01 | 0.95 $\pm$ 0.02 | 0.01     |

#### B. OAR Sparing:

**Table 2. OAR Dose Constraints**

| OAR                    | Parameter       | 10 MV FFF (Mean $\pm$ SD) | 10 MV FF (Mean $\pm$ SD) | p-value      |
|------------------------|-----------------|---------------------------|--------------------------|--------------|
| <b>Rectum</b>          | V50Gy (%)       | 12.5 $\pm$ 3.1            | 14.7 $\pm$ 3.8           | 0.01         |
|                        | V65Gy (%)       | 21.8 $\pm$ 5.2            | 24.3 $\pm$ 6.1           | <b>0.003</b> |
| <b>Bladder</b>         | V65Gy (%)       | 8.7 $\pm$ 4.9             | 10.5 $\pm$ 5.3           | <b>0.02</b>  |
| <b>RT Femoral Head</b> | Mean Dose (cGy) | 1580 $\pm$ 210            | 1680 $\pm$ 240           | <b>0.04</b>  |
|                        | Max Dose (cGy)  | 3354 $\pm$ 450            | 3460 $\pm$ 490           | <b>0.04</b>  |
| <b>LT Femoral Head</b> | Mean Dose (cGy) | 1496 $\pm$ 185            | 1595 $\pm$ 210           | <b>0.03</b>  |
|                        | Max Dose (cGy)  | 3220 $\pm$ 420            | 3320 $\pm$ 440           | <b>0.03</b>  |
| <b>Penile Bulb</b>     | Mean Dose (cGy) | 2472 $\pm$ 1200           | 2600 $\pm$ 1300          | 0.12         |

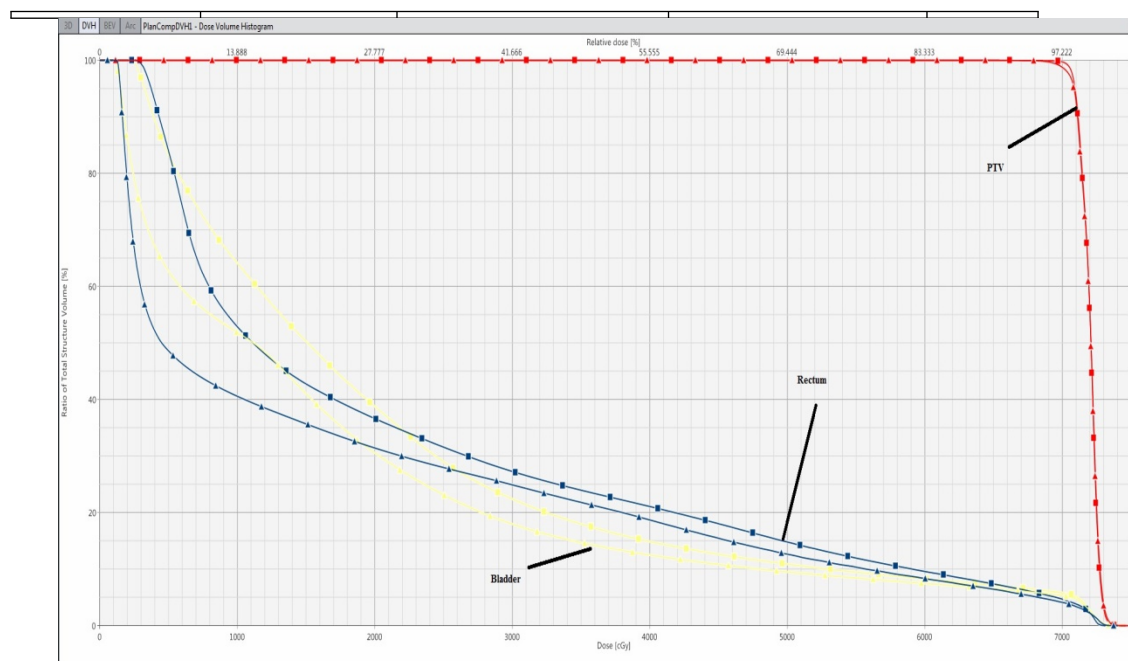


Fig 1.PTV, Bladder & Rectum

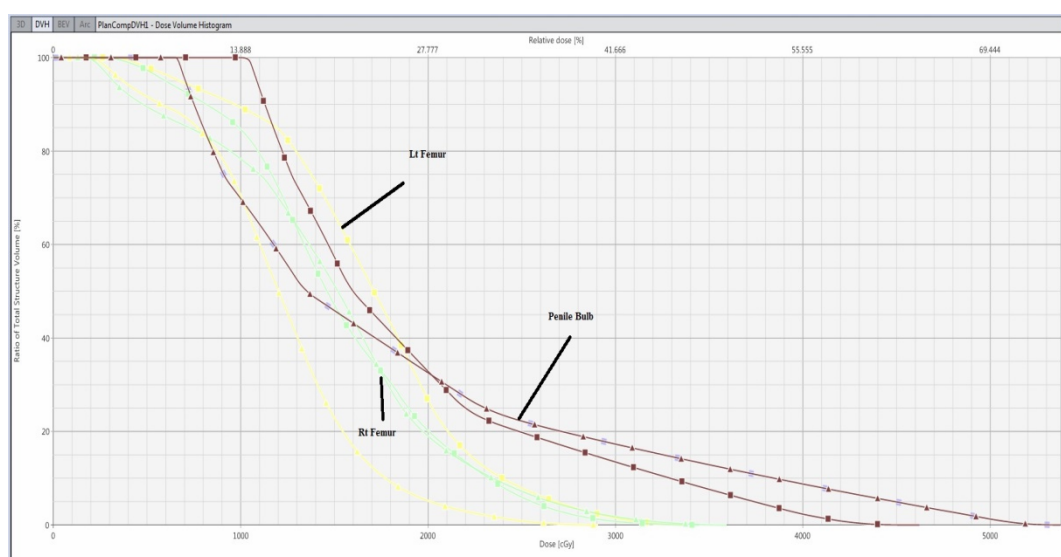


Fig 2.Rt Femur, Lt Femur & Penile Bulb

### C. Treatment Efficiency:

Table 3. Treatment Efficiency

| Parameter            | 10 MV FFF | 10 MV FF  | p-value | Reduction |
|----------------------|-----------|-----------|---------|-----------|
| Treatment Time (min) | 1.7 ± 0.2 | 2.3 ± 0.3 | <0.001  | 26%       |
| Monitor Units        | 880 ± 30  | 1280 ± 50 | <0.001  | 31%       |

### 3. Statistical Analysis

All statistical analyses were performed using paired comparisons between FF and FFF plans for the same patient cohort (n=13). Key methods included:

#### 1. Primary Tests:

- Paired t-tests: For normally distributed data (confirmed by Shapiro-Wilk tests).
- Wilcoxon signed-rank tests: For non-parametric data (e.g., penile bulb doses).

## **2. Multiple Comparisons Correction:**

- Bonferroni adjustment applied to OAR dose metrics ( $\alpha = 0.05/6 = 0.0083$ ).

## **3. Software:**

- IBM SPSS Statistics v28.0 for all analyses.
- Python (SciPy library) for secondary validation.

## **4. Data Presentation:**

- Continuous variables: Mean  $\pm$  Standard Deviation (SD).
- P-values  $<0.05$  considered statistically significant.

## **Discussion**

Our study establishes that 10 MV FFF VMAT is a superior alternative to conventional FF for prostate radiotherapy, with three demonstrable advantages:

### **1. Unmatched Plan Quality:**

- Significant improvements in HI (0.08 vs. 0.11,  $p=0.005$ ) and CI (0.97 vs. 0.95,  $p=0.01$ ) enable sharper dose fall-off near critical structures.

### **2. Enhanced Patient Safety:**

- 10.3% reduction in rectal V65Gy ( $p=0.003$ ) directly addresses one of the most common toxicity concerns in prostate radiotherapy.

### **3. Operational Excellence:**

- The 26% reduction in treatment time (1.7 vs. 2.3 min,  $p<0.001$ ) allows clinics to treat ~8 additional patients daily per machine.

## **IV. Conclusions**

This paired-plan study delivers three pivotal contributions to prostate radiotherapy:

### **1. Technical Advancement:**

The demonstrated superiority of 10 MV FFF VMAT in both plan quality (HI: 0.08 vs. 0.11; CI: 0.97 vs. 0.95) and OAR sparing (rectal V65Gy reduction by 10.3%) establishes it as the new dosimetric benchmark for prostate cancer treatment.

### **2. Clinical Translation:**

- The 26% reduction in treatment time (1.7 vs. 2.3 min) translates to:
- Capacity for 8 additional patients daily per machine.
  - Reduced intrafraction motion risks due to shorter delivery.

### **3. Practical Implementation:**

For clinics with 10 MV linacs, transitioning to FFF requires:

- No hardware modifications.
- Minimal staff retraining (VMAT principles remain unchanged).
- Routine QA adjustments for FFF beam characteristics.

Given these evidence-based advantages, we recommend protocol updates to prioritize 10 MV FFF VMAT for prostate radiotherapy globally.

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