



LOW-RISK ORGAN-CONFINED PROSTATE CANCER



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CyberKnife Center:

Naples Community Hospital Regional Cancer Institute

Naples, FL

LOW-RISK ORGAN-CONFINED PROSTATE CANCER

DEMOGRAPHICS

Sex:

Age: 70

Histology: Prostate Adenocarcinoma: stage T1c

CLINICAL HISTORY

Referred by: Past Medical History: Urologist

Transurethral resection of the prostate (TURP) for benign prostatic hyperplasia (BPH)

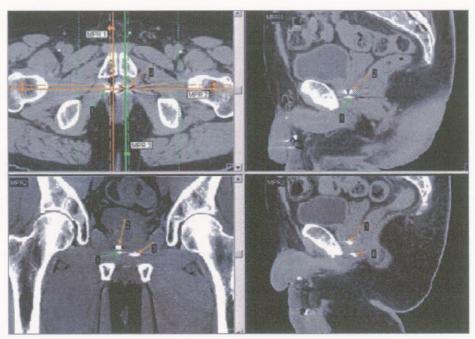
Case History

This 70-year-old male with a history of atrial fibrillation, hypertension and benign prostatic hyperplasia (BPH) presented with elevated prostate specific antigen (PSA) of 4.5 ng/ml in January 2005. He had been followed by his urologist for the previous six years with regular PSA monitoring. He had no family history of prostate cancer and underwent a TURP 2 years prior for BPH. His atrial fibrillation and hypertension were managed by Coumadin, Toprol, Lanoxin and Zestoretic.

The patient's symptoms included nocturia times two and a history of erectile dysfunction. Patient denies a history of dysuria, hematuria, urinary incontinence, urinary urgency, urinary frequency or hesitancy. Transrectal ultrasound (TRUS) guided biopsy revealed adenocarcinoma of the prostate in 6 of 12 biopsy cores, all of which were less than 5% positive and a Gleason score of 3 + 3. Tumor was found in both lobes of the prostate, and was staged cT1c by digital rectal examination. A CT scan of the abdomen / pelvis was unremarkable and a bone scan was negative for metastatic disease.

CyberKnife® Treatment Rationale

The patient was evaluated by Urology and Radiation Oncology for his prostate cancer. Treatment options included surgery, external beam radiation therapy (IMRT, conformal) and CyberKnife monotherapy. The patient wanted a less invasive and convenient therapy in order to continue his work and day to day activities and therefore elected for CyberKnife monotherapy.



Multiplanar pre-treatment planning images show all 4 fiducial markers placed within the prostate.

TREATMENT DETAILS

Prostate Volume: 29.5 cc Imaging Technique(s): CT

Rx Dose & Isodose: 35 Gy to 82%

Conformality Index: 1.39
Tumor Coverage: 95%
Number of Beams: 130

Fractions / Treatment Time:
Path Template:
Tracking Method:

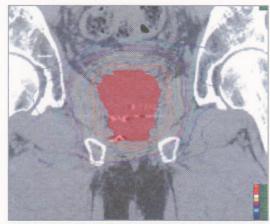
5 @ 43 min per fraction 3 path 900_1000 mm

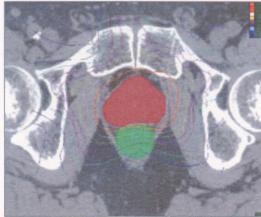
Fiducial

20 mm and 35 mm

Treatment Planning Process

In March 2005, four fiducial markers were placed under intravenous conscious sedation in the prostate by the urologist using a TRUS-guided template. A CT study was performed with the patient in the treatment position using a custom immobilization device. The fiducial locations were identified and the prostate and critical structures (rectum, bladder, and urethra) were contoured. The planning target volume (PTV) included the prostate with a 5-mm margin in all directions except for a smaller 3-mm posterior margin to decrease dosage to the rectum. Treatment planning was designed to encompass 95% of the target volume and minimize dose to critical structures.



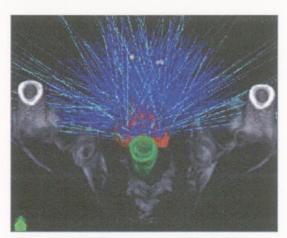


Coronal and axial treatment plans showing the 82% prescription isodose line relative to the prostate (red). Lower percentage isodose lines demonstrate sparing of the rectum (green).

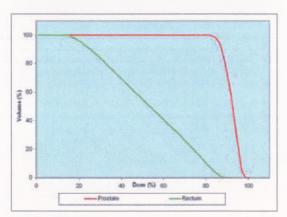
Treatment Delivery

Collimator(s):

The patient began treatment in April 2005. A prescription dose of 35 Gy was delivered in 5 fractions over 5 consecutive days to the 82% isodose line. Two collimator sizes were used and a conformality index of 1.39 was achieved. There were 130 beams from 111 nodes delivered in an average of 43 minutes. Following the fourth treatment, the patient experienced nocturia and was given 0.4 mg Flomax with resolution of symptoms. The patient reported mild urinary frequency and mild urgency 5 days after completion of last fraction of radiosurgery and was treated with Pyridium with resolution of symptoms. Overall, the patient tolerated the treatment well.



Inferior-superior 3D of bony anatomy and CyberKnife beam positions showing treated tumor with rectal sparing.



Dose-Volume Histogram (DVH) for prostate.