

Radiation Therapy After Radical Prostatectomy

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With the current advances in screening, earlier detection of prostate cancer, and improved techniques for performing radical prostatectomy, a larger number of patients are being surgically treated. However, on pathologic examination of their prostate, these patients might not have organ confined disease. The pathologic features which predict failure in those cases are unknown prior to surgery. This includes patients with seminal vesicle invasion, positive surgical margins, and high Gleason scores. Patients at high risk for local recurrence are candidates for further therapy to control the local disease. These patients represent a large proportion of surgically treated patients. In spite of 58% of patients having clinically localized disease at presentation, over one-half will likely be T3 disease.¹ This is despite all the advances in screening.

Adjuvant radiotherapy and/or various methods of androgen deprivation are available for this group of patients. A Mayo Clinic study found that 5-year local recurrence-free survival rates were >95% among prostatectomy patients who had either radiation or orchiectomy compared with 84% for patients who received no adjuvant therapy.² However, no advantage for adjuvant therapy was found in overall or cause-specific mortality. One recent study reported the efficacy of adjuvant radiation after radical prostatectomy, but only among patients found with pT3 who had an undetectable PSA after surgery.³

Postoperative radiation therapy can be given as adjuvant or salvage irradiation. Adjuvant radiation therapy can be used postoperatively for patients with high risk of recurrence. Radiation used as salvage therapy is usually associated with biochemical failure, whether it is associated with a palpable local or biopsy-detected lesion or not. The questions that we need to answer are:

- Does radiation therapy following surgery offer any advantage in local control?
- Does radiation therapy in this setting have any influence on survival and mortality?
- What predicts patients prognosis after radical prostatectomy?
- What should our indications be to use radiation therapy in the postoperative setting?
- How can radiation therapy be used most effectively -- salvage or adjuvant?

EFFICACY OF RADIATION IN THE POSTOPERATIVE SETTING

Men with pathological stage T3 can be further subdivided into prognostically distinct subgroups based on multiple histologic and biochemical factors. Historically, a number of studies with long term follow up

showed that extracapsular disease predisposes local failure. Gibbons et al⁴ had 45 patients with microscopic extension beyond the prostatic capsule. Twenty-two of those patients received external beam irradiation therapy whereas 23 did not. Local recurrence rates (4%) were reduced in the group that received the radiation compared to 30% in the other group that did not. Survival, however, was not affected in either group. Radiation therapy was associated with a higher incidence of pelvic complications. The significance of this finding is of minor importance because of the more advanced techniques of radiation therapy used in recent years. Another study by Shevlin et al⁵ showed that the local recurrence rate can be reduced from 28% to less than 5% with the use of adjuvant radiation.

A more recent study in 1993 by Cheng et al² found that following prostatectomy, the 5 year survival rate was more than 95% in patients receiving radiation or orchiectomy as adjuvant treatment, compared with 84% for other patients not receiving any form of adjuvant therapy. Again, development of distant metastasis and survival were not affected by the adjuvant radiation therapy.

In a recent series,⁶ it was shown that a similar proportion of patients (77%) who received radiation therapy were free of metastatic disease, in comparison with patients (66%) who did not receive radiation. Multiple studies have shown that salvage radiation therapy has variable short term effects on the biochemical status of prostate cancer patients. The probability of developing rising PSA levels after prostatectomy ranges from 18% to 51%, and the duration of undetectable serum PSA is generally short averaging 16 to 24 months in different studies.⁷

Since rising PSA in modern reports is considered the primary indication of failure (local failure or distant metastasis or even both), only those with local failure will benefit from adjuvant radiation therapy. In a recent study by Forman et al,⁸ patients with rising PSA received definitive radiotherapy. Sixty-four percent of patients exhibited no evidence of recurrence up to 48 months after therapy. Treatment failure was 3.7 times greater for patients who did not reach undetectable PSA levels, and 5.6 times greater for patients with either lymph node or seminal vesicle involvement. Patients who had a PSA >1.2 - 2.0 ng/ml postoperatively had a 4 to 5 times greater risk of failing salvage radiation than patients who had PSA values below this mark.

PATTERNS OF FAILURE FOR PT3 PATIENTS

Several studies pointed out that patients with extracapsular disease are predisposed to failure following radical prostatectomy. The local failure rates of Gibbons et al,⁴ Shevlin et al,⁵ and Schellhammer⁹ were 25%, 28% and 31% respectively. Myers and Fleming¹¹ reported a 46% incidence of local recurrence after 10 years. Gibbons et al and Schellhammer suggested that local failure can be predicted by seminal vesicle involvement.

In a multivariate analysis study by Anscher and Prosnitz,¹⁰ the most important predictors of local recurrence were positive surgical margins and high tumor grade. Capsular penetration alone was not a highly suggestive factor. In this same multivariate analysis, seminal vesicle involvement was found to be the strongest predictor of distant metastasis. However, seminal vesicle involvement was not an independent predictor of local failure.¹⁰ The incidence of seminal vesicle involvement with local failure may be higher than reported, but is masked by the fact that patients do not survive due to distant metastasis or the institution of androgen suppression.

Positive Surgical Margins

By strict definition, "positive margins" means neoplastic cells in contact with the inked margin used to mark the specimen. To clarify the positive margin phenomenon, 3 different categories were recognized by Sakr and Grignon:¹²

- Tumor exceeding the inked margin of resection of a seemingly intact anatomical boundary of the gland. In this incidence, local recurrence rates will be similar to organ confined disease.
- Tumor that extends to an irregular, shattered margin indicating surgical induced positive margin. This is commonly a finding at the prostatic apex.
- Tumor extending to definitive extraprostatic fat and/or soft tissue. This is clearly stage T3 cancer.

According to this refined classification by Sakr and Grignon,¹² patients with tumor involving surgical margins but without extraprostatic extension experience biochemical failure rates, in the form of rising PSA, between those of patients with confined tumors and those with extraprostatic disease. Also according to this classification, PSA failure is related to the degree of involvement of the surgical margin.

Patients with positive margins at radical prostatectomy have a 42-64% probability of remaining progression free at 5. This is in contrast to the 80% probability of patients with negative margins remaining progression free at 5 years after radical prostatectomy.^{13,14}

Two studies relating progression to positive margins have followed patients for 10 years. In one of those studies, patients with negative margins had a progression free probability of 74.8%, while those with positive margins had a progression free probability of only 41.9%.¹³ In the other study, progression free probability for patients with positive margins was 38%. Negative margins were not evaluated because these patients were in only 2 separate groups: those with organ confined disease and those with capsular penetration.¹⁵

However, studies found that positive surgical margin was predictive of recurrence only when Gleason sum equal to or greater than 6 with negative seminal vesicle and nodal involvement.¹⁴ In another report, margin status is most relevant in patients with Gleason scores of 5 to 7. In men with Gleason scores 2 to 4, negative seminal vesicles, and negative lymph nodes, prognosis is known to be extremely favorable, while in patients with Gleason scores of 8 or 9, tumors cannot be stratified by capsular penetration or margin status.¹⁶

Huland et al¹⁷ found that in nearly 50% of patients with a Gleason sum of 5 or 6, tumors were organ confined with negative margins. These patients had a 98.7% chance of being disease free at five years, and a 92.4% chance at 10 years, respectively. Since 88% of prostate cancer patients with negative seminal vesicles and nodal involvement fall into this category of a Gleason sum between 5 and 7, the surgical margin status would be of great value in determining the patients' chance of remaining disease free.^{13,16}

Seminal vesicle invasion or lymph node involvement carries a 5 year progression rate of 31% and 23%, respectively. At 10 years, this rises to 58% and 55% respectively. Hence, seminal vesicle invasion or lymph node metastasis patients should be candidates for adjuvant therapy, whether radiation or hormonal. If these findings are detected preoperatively, they should not be candidates for surgery as a means of local control.

IMMEDIATE ADJUVANT OR SALVAGE RADIATION

Historical studies suggest that waiting until gross local recurrence occurs is too late a stage for any local therapy, suggesting it would be more difficult to control gross recurrence versus microscopic residual disease. The Massachusetts General Hospital (MGH) data¹⁸ published in 1993 showed that fewer than 40% of patients irradiated for palpable local recurrences were biochemically disease free at 2 years. The clinical free survival rates were less than 20% at 10 years following radiation therapy. Total "cure" chances are highly unlikely.

However, it is the policy in many institutions to not give radiation to those at risk of local recurrence but rather to wait for signs of biochemical failure. The rationale is that these patients may have no disease or, on the other hand, may have occult metastatic disease. In both cases, they will not benefit from local radiation therapy and will be exposed to the risk of side effects, even though it was reported that patients with occult distant metastasis may briefly benefit from radiation.¹⁹ The results of delayed radiation were also inaccurate due to the fact that some studies included patients with distant metastasis. When lymph

node involvement was excluded, the results were up to 48% freedom from biochemical recurrence at 3 years²⁰ and as high as 68% 3 year freedom from biochemical failure if seminal vesicle invasion was also excluded.²¹ Salvage irradiation also showed some positive results. In contrast to the previous results elsewhere, data from Johns Hopkins Hospital show that 20% of patients were successfully salvaged.²² Nearly half of these patients, however, had seminal vesicle invasion, nodal metastasis, or were at greater risk for distant rather than local recurrence.

The evidence is conflicting regarding the timing of radiation therapy. Long term studies suggest an advantage when adjuvant radiation is given compared to salvage radiation later on. However, this data was obtained prior to the PSA era. Using PSA in follow up means earlier detection of failure. According to Schild et al,²⁰ PSA suppression is inversely related to the PSA level at the time of radiation therapy. This also correlates with the Forman et al data previously discussed in this article. PSA follow up allows us to detect early failure, and hence radiation therapy can be given when tumor volume is small. The critical PSA level was 1.1 ng/ml in the Schild et al study. It is also important that the chance of benefiting from radiation therapy is much higher in delayed rise of PSA beyond the first year of follow up. This means that follow up should be strict and close. If this is not possible or available, early adjuvant radiation may be a reasonable option. Patients who had a PSA >1.2 or 2.0 ng/ml postoperatively had a 4 to 5 times greater risk of failing salvage radiation than patients who had PSA values below this mark.

Radiation's optimal dose varies according to the estimate of tumor burden. The USC group²³ used radiation doses less than those employed by MGH.¹⁸ USC dosage was 45 to 50 Gy versus MGH's 60 to 64 Gy. In the Schild et al study, a dose of 64 Gy was found to be more effective than lower doses, with little morbidity.

CONCLUSION

The main criticism of adjuvant radiation has been its failure to offer survival advantages. This may be due to other competing morbidities masking the survival advantage. Another hypothesis is that the poor biological response of the tumor tends to lead to local recurrence.

The issue of effectiveness of adjuvant radiation still awaits the results of a large Southwest Oncology Group (SWOG) trial. The current SWOG trial examines the benefits of adjuvant XRT with positive margins and capsular penetration after radical prostatectomy. The study randomizes men to adjuvant radiation therapy or to observation. Other data shows the potential benefit with little risk which should not be denied to patients with considerable risk for local recurrence. The patients least likely to benefit from adjuvant or salvage radiation are those with persistently elevated PSA, seminal vesicle invasion, or poorly differentiated tumors. Ultrasound biopsies of the anastomotic site may be helpful to document local recurrence, but negative results should not exclude local disease.

ProstaScint, which is a radiolabeled monoclonal antibody directed towards the prostate specific membrane antigen (PSMA), can be very promising in evaluating patients with rising PSA post radical prostatectomy as well as detection of pelvic lymph node metastasis. PSMA is found in greater amounts in malignant cells and in metastatic deposits, and is also expressed in 95% of all prostate cancer tissues studied. It can very useful in further determining which patients will benefit from post-operative radiation if there is local residual disease.²⁴

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