Scientific Article

Assessment of Polyethylene Glycol Hydrogel Spacer and Its Effect on Rectal Radiation Dose in Prostate Cancer Patients Receiving Proton Beam Radiation Therapy

Anojan Navaratnam, MBBS, FRACS, a Jameson Cumsky, BS, a Haidar Abdul-Muhsin, MB, ChB, a Justin Gagneur, MA, b Jiajian Shen, PhD, b Heidi Kosiorek, MS, c Michael Golafshar, MS, c Akira Kawashima, MD, d William Wong, MD, b Robert Ferrigni, MD, a and Mitchell R. Humphreys, MD a, *

Departments of a Urology, b Radiation Oncology, c Biostatistics, and d Radiology, Mayo Clinic in Arizona, Phoenix, Arizona

Received 10 February 2019; revised 15 July 2019; accepted 13 August 2019

Abstract

Purpose: To assess the efficacy of placing a polyethylene glycol (PEG) spacing hydrogel in patients undergoing proton beam radiation therapy for prostate cancer. This study also aims to assess the effect on rectal radiation dose of prostate—rectum separation in various anatomic planes.

Methods and Materials: Seventy-two consecutive prostate cancer patients undergoing conventionally fractionated pencil beam scanning proton radiation therapy with and without hydrogel placement were compared. Magnetic resonance images taken after hydrogel placement measured prostate—rectum separation and were correlated to rectal dosing and rectal toxicity. Univariate analysis of clinical variables and radiation dosing was conducted using nonparametric Wilcoxon rank-sum test with continuity correction between groups (hydrogel spacer vs controls). Spearman’s rank correlation coefficient assessed relationships between the various anatomic dimensions of perirectal space and rectal radiation dosing.

Results: Fifty-one patients had hydrogel placement before therapy and 21 did not. There was a 42.2% reduction in rectal dosing (mL 3 rectum) in hydrogel patients (P < .001). Increasing midline sagittal lift resulted in a greater mitigation of total rectal dose (P = .031). The degree of prostate surface area coverage on coronal plane did not correlate with further reductions in rectal radiation dose (P = .673). Patients who had PEG hydrogels placed reported more rectal side effects during treatment compared with those patients who did not (35.3% vs 9.5%, P = .061). At median 9.5-month follow-up, there was no difference in reporting of grade ≤2 rectal toxicity between the 2 groups (7.7% vs 7.1%, P = .145).

Conclusions: Polyethylene glycol hydrogel placement before pencil proton beam radiation therapy for prostate cancer reduced rectal radiation dose. The most important factor reducing total rectal dose was the degree of sagittal midline separation created by the PEG hydrogel. This is the largest study with the longest follow-up to investigate hydrogel placement in the proton beam radiation setting.

Sources of support: A.N. received a scholarship from the Australasian Urologic Foundation.

Disclosures: The authors have no conflicts of interest to disclose.

* Corresponding author: Mitchell R. Humphreys, MD; E-mail: humphreys.mitchell@mayo.edu

https://doi.org/10.1016/j.adro.2019.08.007

2452-1094 © 2019 The Author(s). Published by Elsevier Inc. on behalf of American Society for Radiation Oncology. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).
Introduction

Polyethylene glycol (PEG) hydrogel (SpaceOAR, Augmenix, Bedford, MA) is a slowly resorbing hydrogel injected into Denonvillier’s space before external beam radiation (EBRT) for prostate cancer to limit radiation exposure to the rectum. The gel undergoes hydrolysis and dissolves by 6 to 12 months5,6 and was approved by the US Food and Drug Administration (FDA) in 2015.3 Most large series published to date focus on patients undergoing conventional photon beam EBRT2 where utilization of this adjunct has become commonplace.

Displacement of the rectal wall from the prostate by approximately 1 cm allows for a reduction of rectal radiation dose up to 60.6%.1 Recognition of this fact has led to deliberate (saline hydrodissection) efforts to create the space between the prostate and rectum with the hydrogel.

Proton beam radiation therapy (PBRT) is available as an option for men with prostate cancer who choose definitive radiation treatment.4 Proton beams release energy at the Bragg peak, resulting in the benefit of limiting the dose to normal tissue with potential improvement in side effects.4 The rectum has been identified as the dose-limiting collateral structure for traditional photon beam prostate radiation therapy.5 In a randomized trial comparing conventional dose PBRT and high-dose PBRT, Zietman et al reported acute grade 2 or higher gastrointestinal (GI) toxicity as 45% and 64%, respectively.6 Currently, the Prostate Advanced Radiation Technologies Investigating Quality of Life (PARTIQoL) trial (NCT01617161) is ongoing to compare proton therapy to intensity-modulated radiation therapy (IMRT) for low or intermediate risk prostate cancer, with change in bowel function as the primary endpoint. No randomized trials have compared the efficacy and toxicity of PBRT to photon EBRT. Given the proposed advantage of PBRT, the additional benefit of perirectal spacing hydrogel has been questioned. Analysis of Medicare data has previously demonstrated comparatively increased rates of rectal toxicity in PBRT for prostate cancer compared with IMRT during its early adoption,7,8 and hence there is justification for the use of a perirectal spacing agent in this setting.

This is the largest series with the longest follow-up comparing patients with prostate cancer undergoing pencil scanning PBRT with and without placement of PEG hydrogel. The purpose was to determine the effect of the PEG hydrogel on rectal proton beam radiation exposure and relationship between exposure and the degree of prostate—rectum separation. Additional aims were to determine the subsequent effect of PEG hydrogel placement on reducing rectal toxicities from proton beam radiation at early patient follow-up.

Materials and Methods

After institutional review board approval, a retrospective review of patients who underwent conventionally fractionated pencil beam scanning proton EBRT with and without PEG hydrogel placement at Mayo Clinic Arizona was performed.

Patients were positioned in dorsal lithotomy under general anesthesia and 4 carbon fiducial markers were inserted into the prostate. A 15-cm, 18-gauge bevel needle was used to hydro-dissect the potential Denonvillier’s space with minimal normal saline (<2-3 mL) to confirm good needle position before the hydrogel was injected transperineally. A brachytherapy step device was used to hold the rectal ultrasound probe (BK 3000, BK Ultrasound, Peabody MA), and a side-firing biplanar transrectal ultrasound provided real-time visualization ensuring accuracy of hydrogel placement in axial and sagittal planes. A minimum of 1 vial of PEG hydrogel (10 mL) was injected in each patient per manufacturer recommendations.

PBRT was delivered with 2 lateral fields and with 67.5 to 79 Gy in 25 to 44 fractionations depending on the clinical situation. All patients had an endorectal balloon filled with 100 mL of water for each treatment cycle to limit natural rectal displacement.

All patients had a post hydrogel MRI on a 3T platform without an endorectal coil within 7 days to confirm location of the PEG hydrogel and fiducial markers. The following measurements were obtained: (1) Hydrogel thickness: this was measured at the midline of prostate gland on both sagittal and axial images (Fig 1a-b). (2) Maximal surface area of the prostate (AProstate): This was obtained using a reference plane that was 3.51 mm (3 1.17 mm slices on MRI) anterior to the PEG hydrogel (Fig 1c). (3) Maximal contact surface area of the PEG hydrogel (AHydrogel): This was an average of the 2 largest surface area measurements of the PEG hydrogel from coronal images. (4) The difference between points 2 and 3 above and the percentage of prostate coverage were calculated (AOverlap, Fig 2d). Coronal cross-sectional images of the hydrogel and the prostate were superimposed to determine coverage of the prostate using Adobe Photoshop (Adobe Systems, San Jose, CA; Fig 2). Measurements were verified by a single radiologist with expertise in prostate MRI and imaging processing techniques.
The effect of PEG hydrogel placement on rectal radiation exposure was calculated using area under the curve (AUC) for the histogram data of each patient to determine the overall rectal dose based on each patient’s individual dosimetry histogram data (V40 Gy, V50 Gy, V60 Gy, V65 Gy, V70 Gy, and V75 Gy; V indicates the volume of rectum receiving specific radiation dose, eg, V40 Gy is the volume of rectum receiving 40 Gy of radiation). Although previous literature focused on the volume of rectum exposed to the highest recorded radiation dose, the AUC calculation allowed us to estimate overall dosage received by the rectum (Figure E1; available online at https://doi.org/10.1016/j.adro.2019.08.007). Rectal toxicity was graded prospectively by either the clinic nurse or physician at the time of patient follow-up. The follow-up protocol was not standardized and was based on the National Comprehensive Cancer Network guidelines. Toxicities were graded prospectively by the National Cancer Institute Common Terminology Criteria for Adverse Events version 4.03.

Results

Seventy-two consecutive patients with prostate cancer treated with pencil scanning PBRT from January 2016 to August 2017. Fifty-one patients received PEG hydrogel...
before therapy compared with 21 patients who did not. Table 1 details baseline patient characteristics. There were no differences between the 2 groups at median follow-up of 9.5 (8.6-11.5) months after treatment.

Forty-seven (92.2%) patients had 1 hydrogel vial (10 mL) injected, and 4 (7.8%) patients had 2 vials injected. Mean volume of hydrogel injected was 10.75 ± 2.67 mL. All patients had concurrent placement of 4 carbon fiducial markers. No patients experienced any immediate complications.

Median midline separation produced by PEG hydrogel was 10.5 mm (9.4-12.1) and 10.1 mm (8.6-11.4) on

### Table 1  Baseline patient characteristics

<table>
<thead>
<tr>
<th></th>
<th>PEG hydrogel placement (n = 51)</th>
<th>Controls (n = 21)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median age (IQR)</td>
<td>73.9 (70.0-78.0)</td>
<td>74.9 (73.0-78.05)</td>
<td>.291</td>
</tr>
<tr>
<td>Median BMI (IQR)</td>
<td>26.7 (4.8-30.1)</td>
<td>26.3 (24.9-30.0)</td>
<td>.78</td>
</tr>
<tr>
<td>Median PSA (ng/mL) (IQR)</td>
<td>6.9 (4.5-10.2)</td>
<td>9.7 (4.8-12.4)</td>
<td>.222</td>
</tr>
<tr>
<td>Patient on ADT</td>
<td>26 (n = 40, 65%)</td>
<td>19 (n = 10, 95%)</td>
<td>.027</td>
</tr>
<tr>
<td>Gleason score</td>
<td></td>
<td></td>
<td>.361</td>
</tr>
<tr>
<td>3 + 3</td>
<td>10 (19.6%)</td>
<td>1 (4.8%)</td>
<td></td>
</tr>
<tr>
<td>3 + 4</td>
<td>15 (29.4%)</td>
<td>4 (19.0%)</td>
<td></td>
</tr>
<tr>
<td>4 + 3</td>
<td>15 (29.4%)</td>
<td>8 (38.1%)</td>
<td></td>
</tr>
<tr>
<td>4 + 4</td>
<td>7 (13.7%)</td>
<td>6 (28.6%)</td>
<td></td>
</tr>
<tr>
<td>4 + 5</td>
<td>2 (3.9%)</td>
<td>2 (9.5%)</td>
<td></td>
</tr>
<tr>
<td>5 + 4</td>
<td>1 (2.0%)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>5 + 5</td>
<td>1 (2.0%)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Clinical stage</td>
<td></td>
<td></td>
<td>.65</td>
</tr>
<tr>
<td>T1 (T1a, T1b, T2c)</td>
<td>22 (43.1%)</td>
<td>8 (38.1%)</td>
<td></td>
</tr>
<tr>
<td>T2</td>
<td>24 (47.1%)</td>
<td>12 (57.1%)</td>
<td></td>
</tr>
<tr>
<td>T3 (T3a, T3b)</td>
<td>5 (9.8%)</td>
<td>1 (4.8%)</td>
<td></td>
</tr>
<tr>
<td>Median radiation dose delivered Gy (IQR)</td>
<td>79.2 (79.2-79.2)</td>
<td>79.2 (79.2-79.2)</td>
<td>.621</td>
</tr>
<tr>
<td>Median no. of fractions delivered</td>
<td>44.0 (44.0-44.0)</td>
<td>44.0 (44.0-44.0)</td>
<td>.786</td>
</tr>
</tbody>
</table>

Abbreviations: ADT = androgen deprivation therapy; BMI = body mass index; IQR = interquartile ratio; PEG = polyethylene glycol; PSA = prostate-specific antigen.
sagittal and axial planes, respectively. Only 32 patients had the necessary coronal images to calculate coverage of prostate by the PEG hydrogel in the coronal plane. Median percentage of prostate covered by the PEG hydrogel in coronal plane ($A_{\text{overlap}}$) was 49.0% (39.9-57.7).

Patients who had PEG hydrogel placement had reduced radiation dose per volume of rectum at all histogram dosimetry levels compared with those with no PEG hydrogel ($P < .001$; Figure E2; available online at https://doi.org/10.1016/j.adro.2019.08.007). Volume of rectum exposed was reduced by 42.2% and 41.8% for V70 Gy and V75 Gy, respectively. AUC analysis also demonstrated a 42.2% relative reduction in overall rectal dose delivery in hydrogel patients ($P < .001$; Fig 3).

When overall rectal dosimetry based on AUC calculation was compared with prostate–rectum separation measurements, greater mitigation of rectal dosing was seen with increased midline prostate–rectum separation on sagittal MRI ($P = .031$). Increased coverage of the prostate by PEG hydrogel in the coronal plane was not associated with reduced rectal radiation dose based on our measurements (Fig 4).

Patients with hydrogel placement reported more toxicity during treatment compared with controls. At least one grade 1 acute rectal toxicity was experienced in 35.3% of hydrogel patients compared with 9.5% of controls ($P = .061$) during the course of therapy.

With longer follow-up (median 10.3 vs 8.7 months, hydrogel vs control), the overall reporting of rectal toxicity was not different between the 2 groups (Table 2). There was no association between rectal toxicity at any point and PEG hydrogel measurements.

**Discussion**

The success of PEG hydrogel spacers in significantly reducing short- to medium-term rectal toxicity has been well documented in the photon EBRT setting. In this pivotal trial that led to the approval of hydrogel by the FDA, no rectal balloon was used. Placement of hydrogel is well tolerated, improves GI quality of life scores at 5 years, and has been shown to be cost effective. PBRT alone has previously demonstrated low rates of rectal toxicity even without the use of a spacer. The added benefit of placing a PEG hydrogel spacer in this setting has not been well investigated and can be questioned given the theorized reduction in collateral radiation from proton beam radiation. Only a single small series of 12 patients has previously demonstrated a relative reduction in V70 Gy rectal radiation dose of 63% and 70% based on 2 separate proton radiation plans using an alternative PEG Hydrogel (Duraeseal, Covidien, Mansfield, MA). We report the largest study to directly compare PBRT patients treated with and without PEG hydrogel placement.

Our study demonstrated a 42.2% relative reduction of rectal exposure in PBRT patients with the use of the hydrogel spacer during treatment. Each rectal dosimetry level demonstrated significantly less radiation exposure. This is consistent with previously published results in the photon EBRT population, which have demonstrated a relative rectal dose reduction of 25% to 59% in patients who had PEG Hydrogel placed before therapy.

A median prostate–rectum separation at midline of 10.5 mm on sagittal imaging was achieved in the hydrogel group and was positively correlated with dose reduction. A statistically significant association was not seen between dose reduction and length of separation on axial images. Pinkawa et al have previously demonstrated that there was a learning curve required to obtain up to 15 mm of separation from of the prostate to the anterior rectal wall. Preclinical studies have shown that no further reduction in V70 Gy occurred with separation $>15$ mm after 20 mL of hydrogel injection. Furthermore, these results are consistent with data in the photon EBRT population demonstrating separation of 10 to 13 mm is most effective at reducing rectal exposure. Cadaver and small clinical series of treatment planning with PBRT did demonstrate that 7 to 9 mm of prostate–rectum separation can result in reduced rectal radiation dose.

Coronal coverage maps demonstrated that a median of 49.0% of the contact surface area of the prostate was covered by the PEG hydrogel. Often, $<50\%$ of the contact surface of the prostate was overlapped by the PEG hydrogel in the coronal plane due to displacement of the hydrogel away from the midline. This suggests that the hydrodissection of the potential space of Denonvillier’s may not be predictable or that hydrogel...
placement is more unpredictable than wanted. It was anticipated that the degree of prostate covered by the hydrogel on coronal imaging would correlate with a more uniform separation of the prostate from the rectum and greater rectal dose reduction; our study did not demonstrate this association. This questions the necessity of ensuring even distribution of hydrogel in the peri-rectal space rather than focusing on the goal of adequate lift of the prostate at midline.

No relationship was found between the thickness of prostate—rectum separation at midline on imaging and rectal toxicity reported either during treatment or on follow-up. Although increasing thickness of separation may improve dosimetry scores, its clinical effect remains uncertain. Mariados et al demonstrated that hydrogel absorption began at 3 months and completely resolved by 12 months. It is difficult to assess the clinical effect of initial separation given the variable rates of absorption occurring between patients owing to differences in renal clearance and proteolytic resistance.

During treatment, greater rectal toxicity was reported in patients with PEG hydrogel placement than without. It must be acknowledged that this is a near-significant approximate 4-fold increase of in-treatment rectal toxicity in those patients who received PEG hydrogel. None of these patients needed intervention for these side

![Figure 4](image)

**Figure 4** Comparison of Hydrogel placement with area under the curve (AUC) rectal radiation dosage (cc3.Gy) with Spearman rank correlation calculation (rho). (a) Sagittal thickness of hydrogel in midline (cm) versus AUC ($P = .031$). (b) Axial thickness of hydrogel in midline (cm) versus AUC ($P = .222$). (c) Percentage of prostate coverage/overlap (AOverlap) in coronal plane versus AUC ($P = .673$).
effects and all toxicities were grade 1 and did not require any intervention. This increased propensity for hydrogel patients to experience rectal side effects may be due to interaction between the rectal balloon and PEG hydrogel in place concurrently with each radiation treatment (Fig 1a-b demonstrating proximity of the 2 structures). Rectal balloons have been used in PBRT to reduce variability in prostate position given the increased sensitivity of PBRT to target motion owing to steep dose depletion beyond the Bragg peak. We have used an endorectal balloon for PBRT patients as a method of reducing intrafraction movement, extrapolating from previous studies of IMRT for prostate cancer. Small studies have described patients complaining about local side effects from PEG hydrogel placement. A recent retrospective review of 125 patients undergoing photon EBRT revealed an increased rate of hemorrhoids in patients who had hydrogel placement. Another study described transient increase in rectal discomfort in 4 of 11 patients who received PEG hydrogel before photon EBRT. These side effects resolved within 12 weeks of hydrogel placement.

Interaction of the PEG hydrogel and rectal balloon on the anterior rectal wall may represent a side effect of the hydrogel as opposed to true radiation toxicity. It is also possible that Common Terminology Criteria for Adverse Events, version 4 is not sensitive enough to differentiate rectal discomfort secondary to the simultaneous use of the hydrogel and rectal balloon, and the discomfort was graded as proctitis at the weekly management visit during the treatment course. The concurrent placement of a PEG hydrogel and endorectal balloon requires more investigation of its effect on intrafraction prostate movement.

Fifty-three patients (73.6%) had completed follow-up at a median time of 9.5 (8.6-11.5) months. Of these, only 7.5% experienced grade ≤2 rectal toxicity. There was no difference in rectal toxicity between the 2 groups at early follow-up. The median follow-up time for the hydrogel group was 1.6 months longer than the control group. A possible explanation is that the physical discomfort caused by the hydrogel and rectal balloon resolved after resorption of the hydrogel in 3 to 6 months. Large series of PBRT patients without hydrogel have variable reported rectal toxicity rates ranging from 0% to 64%. The pivotal randomized controlled trial that led to FDA approval of PEG hydrogel in photon EBRT demonstrated no difference in acute toxicity rates (<3 months after therapy) with 23% and 28% in hydrogel and control groups, respectively. However, at late follow-up (>3 months), there was a significant reduction in rectal toxicity (grade ≤2) in those who had hydrogel placement of 7% versus 2%. This study demonstrates no difference in rectal toxicity rates on longer follow-up (>3 months). This was recently updated with 3-year follow-up and it was demonstrated that patients with PEG hydrogel experienced statistically significantly less rectal toxicity compared with controls. They also demonstrated that improved bowel quality of life scores (Expanded Prostate Cancer Index Composite) were maintained at 3 years in patients who had hydrogel placement. The authors determined that the number needed to treat to spare grade ≥1 and ≥2 toxicity at 3 years were 1.3 and 16.7, respectively, which raises the clinical significance of these improvements. Our study requires longer term follow-up to determine the effect of the decrease in rectal dose secondary to PEG hydrogel placement on reporting of rectal toxicity in PBRT. Although this study also investigated the effect of PEG hydrogel on urinary toxicity, this was not an endpoint of our study.

There were several limitations in this study. The measured height of prostate—rectum separation at a single midline point in the sagittal and axial planes was arbitrarily determined. Although this allowed us to measure the length of separation in a standardized fashion, measurements at other points of reference may have resulted in different correlations between length of separation and rectal radiation dose. The study was also limited by its relatively small number of patients and may be underpowered to detect differences between the groups. Additionally, because both PEG hydrogel and pencil beam scanning PBRT are relatively new therapies for prostate cancer treatment, the patients in our study have a relatively short follow-up. Additional follow-up is required to compare the effectiveness of PBRT with
hydrogel spacer to photon EBRT with hydrogel spacer in reducing long term rectal toxicity. Finally, the retrospective nature of our review resulted in inconsistencies of data reporting, particularly pertaining to toxicity scores.

Conclusions

This is the largest study to investigate the effect of PEG hydrogel placement in patients undergoing proton beam radiation therapy for prostate cancer. PEG hydrogel placement before pencil beam scanning PBRT reduces overall rectal dose by 42.2%. Increased midline separation in the sagittal plane correlates with reduced rectal dosing. Placement of the hydrogel in Denovillier's space may not be predictable given the relatively low overlap between hydrogel and prostate on coronal imaging; however, this does not affect the efficacy of the PEG hydrogel. Concurrent presence of the PEG hydrogel and rectal balloon during PBRT may result in increased reporting of rectal toxicity during treatment, which may represent a shortfall of the reporting system to differentiate rectal discomfort from true radiation toxicity. There was no difference between the rates of rectal toxicity reported between patients with and without PEG hydrogel at 9.5 months after completion of PBRT. Additional follow-up will allow the determination if placement of PEG hydrogel in the PBRT setting results in reduction in long-term rectal toxicity, as well as ensuring there is no deterioration in the treatment effect of PBRT.

Supplementary data

Supplementary material for this article can be found at https://doi.org/10.1016/j.adro.2019.08.007.

References


