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The dosimetric impact of vaginal balloon-packing on intracavitary high-dose-rate brachytherapy for gynecological cancer

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Abstract

**Purpose:** We perform a clinical retrospective study to determine whether a vaginal balloon-packing system provides a dosimetric reduction to organs at risk (OARs) versus traditional gauze packing for gynecological high-dose-rate brachytherapy (HDR-BT). We also test various balloon filling materials for optimizing imaging quality.

**Material and methods:** Filling materials for balloon-packing were evaluated based on imaging quality with X-ray, computerized tomography, and magnetic resonance imaging modalities. We then retrospectively reviewed 45 HDR-BT plans of 18 patients performed with gauze packing and 39 plans of 16 patients performed with balloon-packing. Twelve patients received both gauze and balloon-packing. HDR-BT was delivered with an iridium-192 afterloader and a Fletcher-Suit-Declos-style T&O applicator. At each fraction, 3D imaging was obtained. The D2 cc values of OARs were calculated, as well as ICRU-defined point doses.

**Results:** In the 84 HDR fractions reviewed, vaginal balloon-packing provides statistically equivalent doses to rectum, bladder, and sigmoid compared to gauze packing. On average balloon-packing produced average reductions of 3.3% and 6.9% in the rectal and sigmoid D2 cc doses and an increase of 3.2% to the bladder D2 cc dose (normalized to prescription dose), although none of these values were statistically significant for the twelve patients who received both gauze and balloon-packing (32 and 40 total fractions, respectively).

**Conclusions:** In the 84 HDR fractions analyzed, vaginal balloon-packing is as effective as gauze packing for dose sparing to the rectum, bladder, and sigmoid. A 1 : 1 solution of saline and contrast for filling material enables easy contouring for image-guided HDR with minimal artefacts.

Key words: brachytherapy, gynecological cancer, high-dose-rate brachytherapy, vaginal balloon packing.
in HDR-BT is currently gaining momentum [8,9] due to its high soft tissue contrast, which provides better definition of the macroscopic tumor. Commonly referenced volumetric dose parameters include the $D_{2cc}$, which is the dose in the “hottest” 2 cc of a structure, as opposed to the traditional rectal and bladder point doses defined on 2-D imaging by the International Commission on Radiation Units and Measurements (ICRU) report #38 [5].

While a previously published study used Monte Carlo modelling to show a theoretical 6-15% dose reduction to OARs using vaginal balloon-packing [10] due to the attenuating effect of a high CT-number contrast material used as filling, to our knowledge, no clinical studies concerning the dosimetry of vaginal balloon-packing have been reported. In this study, we investigate whether balloon-packing provides superior dose reduction to OARs compared with traditional gauze packing. We hypothesize that the increased reproducibility afforded by the balloon-packing system and improved treatment geometry would on average lead to decreased OAR doses compared to gauze packing, due to the inherent variability in quality of gauze packing. Toward this end, we describe a retrospective volume-based dosimetric analysis of OAR doses from HDR-BT plans performed with either gauze or balloon-packing. Additionally, dosimetric comparisons are made among fractions for patients who received both gauze and balloon-packing during the course of their treatment.

Material and methods

Patients and treatment: EBRT and HDR

Twenty-two patients with biopsy-confirmed cervical cancer who received HDR-BT were selected for this study. The use of patient data was approved by our Institutional Review Board. The patients’ International Federation of Gynecology and Obstetrics (FIGO) stages ranged from IB2 to IIIB. Institutional standard-of-care treatment consisted of external beam radiation therapy (EBRT) with weekly chemotherapy (typically platinum based) integrated with HDR-BT. EBRT was 45 Gy in 25 fractions to the whole pelvis. Patients with parametrial involvement received an EBRT boost of 5.4 Gy in 3 fractions to the parametrium. After 3-4 weeks of EBRT, HDR-BT using a Varian Medical System Inc., Palo Alto, California, USA) was started with a nominal 7 day interval, integrated with EBRT so that the radiation course is finished within 7 weeks [3]. HDR treatment plans were performed based upon the conventional Manchester-based system [3,13], in which radiation dose is prescribed to point A. The BrachyVision version 8.9® (Varian Medical Systems Inc.) treatment planning system was used for HDR planning and dose volume histogram analysis.

Vaginal packing procedure

After the T&O applicator was positioned, either gauze or balloon-packing (the Alatus® Vaginal Balloon Packing System, Radiadyne LLC., Houston, Texas, USA) was used to immobilize the applicator. For gauze-packing, a roll of gauze was soaked in sterile normal saline. Beginning at the proximal vagina, approximately 5 cm segments of gauze were inserted with forceps, alternating anteriorly and posteriorly to the applicator. Gauze insertion continued until the vaginal introitus was reached and the applicator remained snugly in place.

The balloon-packing system consists of two balloons which are inserted anteriorly and posteriorly to the applicator. A 50 cc syringe was filled with a solution of one part normal saline and one part ISOVUE-250® contrast (Bracco Diagnostics Inc., Princeton, New Jersey, USA). This syringe was used to alternately fill the anterior and posterior balloons in 10 cc increments. Each balloon was typically filled with 30 cc to 50 cc of solution depending on the vaginal size.

Commissioning vaginal balloon-packing

Various balloon filling materials were examined, including air, water, 100% contrast, 100% normal saline, and a 1:1 solution of normal saline and contrast (Fig. 1). The optimal filling material was determined using two metrics: favourable imaging qualities on orthogonal radiograph, CT, and MRI modalities (most importantly lack of artefact), and also radiation attenuation capacity. A quality assurance (QA) phantom, previously developed to perform QA for CT or MRI-guided HDR [11], was also used in this study. The Hounsfield Unit (HU) values on CT were quantified to estimate the radiation attenuation potential for the various filling materials. The vaginal balloon-packing material which best balanced imaging characteristics with the potential for dose attenuation was clinically implemented.

Dosimetric evaluation

To perform in vivo dosimetric evaluations, we retrospectively reviewed 45 HDR plans of 18 patients that were performed with gauze packing and 39 plans of 16 patients performed with balloon-packing. Twelve patients received both gauze and balloon-packing. The rectum, bladder, and sigmoid were contoured on high-resolution (3.0 Tesla) MRI
Fig. 1. The first row shows CT images with vaginal balloon-packing with air (A), water (B), 100% contrast (C: anterior balloon), 100% saline (C: posterior balloon), and 1:1 saline-contrast solution (D). Panel (E) shows orthogonal radiographs with water filling and the QA phantom. The second row demonstrates in vivo radiographs (F & G) and (H) T2 & (I) T1-weighted MRI. Panel (J) depicts T2-weighted MRI with conventional gauze-packing.
or CT scans, along with the GEC-ESTRO-EMBRACE guidelines [8]. Values for D$_{2cc}$ of the OARs were used as metrics to compare the dosimetric impact of vaginal packing, along with conventional ICRU report #38-defined rectal and bladder point doses. Both the ICRU point doses and the D$_{2cc}$ values were converted to a percentage-of-prescription dose for that fraction by dividing by the prescription dose and multiplying by 100 (referred to as %ICRU or %D$_{2cc}$). This was done to normalize to the prescription dose (5.5 Gy or 7 Gy). The average ICRU, %ICRU, D$_{2cc}$, and %D$_{2cc}$ values for the OARs were calculated for gauze and balloon-packing, as well as standard deviations. The $p$-values were calculated using Microsoft Excel using a two-tailed $t$-test assuming homoscedasticity.

An “intra-patient” comparison of balloon and gauze packing was conducted for twelve patients who received both gauze and balloon-packing at different fractions during the course of their treatments. These patients each received 1-3 fractions packed with gauze and 1-4 fractions packed with balloons. The %ICRU and %D$_{2cc}$ doses were calculated as described above for the ICRU point doses and the image-based D$_{2cc}$ values, respectively. For those patients who received more than one fraction with a particular packing type, values were averaged and the standard deviation was calculated.

Results

Balloon filling material

We found that air filling was suboptimal for balloon-packing due to the loss of attenuation potential. Saline provided both a degree of attenuation (62 ± 32 HU) and satisfactory image quality on radiographs and CT, showing the vaginal wall clearly with no artefacts, and on T2-weighted MRI with high MR signal. Contrast used as filling produced a high HU (2790 ± 585), which was potentially promising for further attenuation to the OARs, but imaging artefacts were unacceptable (Fig. 1). A 1 : 1 mixture of saline and contrast produced acceptable imaging on orthogonal radiographs, CT, and MRI with a HU value of 1882 ± 449, also promising for potential attenuation. The 1 : 1 saline and contrast filling was selected for our institutional protocol for MRI-guided BT. Both radiographs and MRI (T1 & T2-weighted) with balloon-packing showed the vaginal wall clearly, aiding with contouring during treatment planning.

Dosimetric analysis

A summary of the dosimetric comparison between balloon-packing and gauze packing is shown in Table 1. Balloon-packing provided a non-statistically significant reduction in the D$_{2cc}$ rectal dose expressed as a percentage of the prescription (53.1% balloon vs. 60.7% gauze, $p = 0.07$). Likewise, the bladder and sigmoid D$_{2cc}$ volumetric doses and the ICRU rectal and bladder point doses were statistically equivalent between balloon-packing and gauze packing. The bladder D$_{2cc}$ dose was higher for balloon packing (98.1% balloon vs. 87.7% gauze, $p = 0.16$), but again, this was not statistically significant.

In order to account for anatomic differences, a comparison of OAR doses was made among fractions within the same patients, i.e., 12 patients who received both gauze and balloon-packing alternatively at different fractions during the courses of their treatments (Table 2). Each patient received 1-4 fractions of both packing types. For these twelve patients as a group, balloon-packing produced average reductions of –3.3%, and –6.9% in the rectal and sigmoid D$_{2cc}$ doses, and an increase of +3.2% in the bladder D$_{2cc}$ dose (expressed as a percentage of the prescription dose). There were -3.5% and –4.8% reductions in the rectal and bladder ICRU point doses as a percentage of the prescription doses. None of these values were statistically significant.

Discussion

Gauze has been utilized as vaginal packing for gynecological BT for decades with traditional 2-D orthogonal imaging. The quality of gauze packing is highly dependent on the experience and skill of the oncologist performing the procedure. Potential pitfalls include not packing tightly enough or an uneven distribution of gauze between the anterior and posterior parts of the applicator. At our institution, the applicator is placed while the patient is under general anaesthesia, but aggressive packing can be very uncomfortable for sedated patients. The vaginal wall is also susceptible to laceration during the packing process, causing further potential morbidity to the patient and increasing the total procedure time if sutures are required.

Balloon-packing addresses many of the problems with gauze packing. We hypothesized that balloon-packing would decrease dose to the OARs by increasing distance from sources and a more favourable geometry. Saini et al. [10] reported contrast-filled balloons showed 7.8% and 19.2% dose reductions through diode measurement when compared to saline- and air-filled balloons, respectively. Monte Carlo modelling also predicted 10.5% and 21.9% dose reductions in contrast-filled balloons, compared to saline- and air-filled balloons, respectively. In this study, ISOVUE-250 contrast was used in which iodine is the effective element and has 33 keV K-shell electron binding energy. Thus, iodine has a high attenuation coefficient due to the photoelectric effect for low energy X-rays or gamma-rays. Enhanced photon absorption in the solution is, therefore, expected to be due to the large number of the photoelectric events, because of the presence of iodine. No study has been reported, validating dose reduction on OAR due to the use of contrast-filled balloon packing using clinical datasets. This is an area for future research.

In our analysis of 84 HDR fractions, we did not find that balloon packing made a statistically significant difference in the bladder, rectal, or sigmoid OAR doses using either ICRU point criteria or a volumetric D$_{2cc}$ analysis. Anatomical variation in these OARs may explain why there was not a significant reduction in doses to these structures with balloon-packing compared to gauze packing. This variation is seen among patients and also among fractions for the same patient. In an attempt to account for the former, we compared both packing types in twelve patients who received gauze and balloon-packing during the courses of their treatments. In this analysis, balloon-packing did produce a slight average reduction in dose for the rectum and sigmoid (3.3%...
Table 1. Retrospective dosimetric analysis of 45 HDR brachytherapy plans performed with gauze packing and 39 plans performed with vaginal balloon packing. Mean values and standard deviations are shown below, as well as p-values from a two-tailed t-test. The average dose per fraction was similar between the gauze packing and balloon packing groups (6.1 Gy and 6.2 Gy). ICRU rectal and bladder point doses and the D2cc values for the rectum, bladder, and sigmoid were calculated. %ICRU and %D2cc refer to the ICRU and D2cc values as a percentage of the prescription doses.

<table>
<thead>
<tr>
<th>Prescription</th>
<th>Rectum</th>
<th>Bladder</th>
<th>Sigmoid</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Gauze</td>
<td>Balloon</td>
<td>Δ</td>
</tr>
<tr>
<td>ICRU</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3.7 ± 1.3 Gy</td>
<td>6.1 ± 0.7 Gy</td>
<td>6.2 ± 0.7 Gy</td>
<td>0.52</td>
</tr>
<tr>
<td>%ICRU</td>
<td>34.7 ± 19%</td>
<td>53.1 ± 18.4%</td>
<td>−7.6%</td>
</tr>
<tr>
<td>D2cc</td>
<td>3.8 ± 11 Gy</td>
<td>3.5 ± 1.2 Gy</td>
<td>−0.3 Gy</td>
</tr>
<tr>
<td>%D2cc</td>
<td>62.4 ± 14.2%</td>
<td>56.5 ± 17%</td>
<td>−5.9%</td>
</tr>
</tbody>
</table>

Prescription – prescriptions for all 45 HDR plans, Gauze – gauze packing data, Balloon – vagina balloon packing data, Δ – the mean difference of balloon-gauze (negative represents dose reduction at balloon), p – p values, ICRU – ICRU report #38 defined rectal and bladder point doses, %ICRU and %D2cc – ICRU and D2cc values as a percentage of the prescription doses.

Table 2. The dosimetric differences of 12 intra-patients who received both gauze and balloon packing. Negative represents dose reduction in balloon packing.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Difference in rectum</th>
<th>Difference in bladder</th>
<th>Difference in sigmoid</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>ICRU [Gy] %ICRU [%]</td>
<td>D2cc [Gy] %D2cc [%]</td>
<td>ICRU [Gy] %ICRU [%]</td>
</tr>
<tr>
<td>1</td>
<td>−11</td>
<td>−15.2</td>
<td>−3.0</td>
</tr>
<tr>
<td>2</td>
<td>+0.7</td>
<td>+12.7</td>
<td>+0.3</td>
</tr>
<tr>
<td>3</td>
<td>−0.2</td>
<td>−3.1</td>
<td>+1.0</td>
</tr>
<tr>
<td>4</td>
<td>−3.1</td>
<td>−43.8</td>
<td>−2.2</td>
</tr>
<tr>
<td>5</td>
<td>−1.3</td>
<td>−23.5</td>
<td>−0.7</td>
</tr>
<tr>
<td>6</td>
<td>+0.2</td>
<td>+6.6</td>
<td>−0.9</td>
</tr>
<tr>
<td>7</td>
<td>+15</td>
<td>+20.7</td>
<td>+2.0</td>
</tr>
<tr>
<td>8</td>
<td>+0.5</td>
<td>+7.1</td>
<td>+0.7</td>
</tr>
<tr>
<td>9</td>
<td>−0.1</td>
<td>−1.0</td>
<td>−0.2</td>
</tr>
<tr>
<td>10</td>
<td>+0.3</td>
<td>+4.0</td>
<td>0.0</td>
</tr>
<tr>
<td>11</td>
<td>−0.9</td>
<td>−16.0</td>
<td>+0.3</td>
</tr>
<tr>
<td>12</td>
<td>+0.5</td>
<td>+9.4</td>
<td>−0.3</td>
</tr>
<tr>
<td>Average</td>
<td>−0.2</td>
<td>−3.5</td>
<td>−0.2</td>
</tr>
<tr>
<td>Std dev</td>
<td>1.2</td>
<td>19.1</td>
<td>1.5</td>
</tr>
<tr>
<td>P value</td>
<td>0.52</td>
<td>0.48</td>
<td>0.58</td>
</tr>
</tbody>
</table>

Difference – the difference of balloon packing data – gauze packing data, ICRU – ICRU report #38 defined rectal and bladder point doses, %ICRU and %D2cc – ICRU and D2cc values as a percentage of the prescription doses.

and 6.9%), but an equally slight increase in dose to the bladder (3.2%). These values were not statistically significant.

Among these twelve patients, many patients often had suboptimal high bladder doses for at least one fraction, regardless of the packing type (Table 2). Notably, the ICRU bladder point dose was often deceptively within normal limits (< 80% of the prescribed dose), but the image-based D2cc was over 100% of the prescribed dose. The disconnection between ICRU point doses and image-based dosimetry has been previously noted [14], and image-based dosimetry has been shown to be more clinically relevant in terms of side effects [15]. Our current study further underscores the importance of volume-based dosimetry in brachytherapy planning.

In a patient with a very anteverted uterus, portions of the bladder may be unavoidably near the tandem and therefore in the high-dose radiation isodose levels regardless of the type of packing used. Similarly, the anatomic location of the sigmoid colon varies among patients and among fractions, and its location with respect to the radiation isodose distribution is often unaffected by the type or degree of packing, since it is physically distant from the vagina.

When we started using balloon-packing in our practice, we filled both the anterior and posterior balloons to the no-
minal 40 cc as recommended by the vendor. Most patients in this study were treated according to these instructions. In contrast, when using gauze packing, the volume of packing is not set for each fraction, and the amount of packing a patient receives may vary from fraction to fraction based on the physician’s skill and judgment. Nevertheless, we demonstrate that even with vendor recommended filling, we achieve, on average, comparable, if not better, dose sparing to OARs with balloon-packing. We believe that this demonstrates the major benefit of balloon-packing, which is a consistent quality of packing from fraction to fraction that minimizes the reliance on technique. We have recently begun to increase the filling volume of each balloon to 50 cc if allowed by the patient’s anatomy, which could conceivably provide some additional degree of dosimetric sparing to the OARs.

We show in this study that while there is not a dosimetric advantage to using balloon packing, an extensive analysis of 84 HDR fractions shows balloon packing provides statistically equivalent dosimetric protection to the OARs. This should provide reassurance to clinicians using balloon packing for reasons of patient comfort and convenience and may also suggest balloon packing as an appropriate selection for low-volume HDR centers with less experience with gauze packing. There are not enough patients and there has not been sufficient follow-up to know whether there are any differences in clinical outcomes between the two groups of patients. In the short term, we have not noted any significant differences in the acute side effect profiles.

Conclusions

To our knowledge, this is the first clinical analysis of the dosimetric performance of a commercially available vaginal balloon-packing system. We show that based on volumetric dosimetry, balloon-packing provides a statistically equivalent OAR protection in the 84 HDR fractions analyzed. Balloon-packing provides significant clinical benefits, including patient comfort and safety and time savings. Although we did not show that balloon packing is able to significantly lower OAR doses through geometry optimization, clinical demonstration of a possible dose reduction secondary to attenuation by a high-Z balloon filling material does remain an area for future research.

Despite the promise of using image-guided HDR to deliver more personalized, conformal radiation to patient’s actual tumor volumes as opposed to the historical Manchester points, HDR delivery is ultimately constrained by the physical geometry of the implant, which in turn is dictated by the patient’s anatomy and OAR constraints. New paradigms, including balloon-packing, offer additional tools for increasing the therapeutic ratio of brachytherapy, either by optimizing the geometry of the implant, or possibly through physical means such as introducing some degree of radiation attenuation in the balloon filling material.

Acknowledgements

This work was partially supported by research grants from Varian Medical Systems.

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