Dosimetric comparison of three different treatment modalities for total scalp irradiation: the conventional lateral photon–electron technique, helical tomotherapy, and volumetric-modulated arc therapy

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ABSTRACT

The aim of this study was to compare lateral photon–electron (LPE), helical tomotherapy (HT), and volumetric-modulated arc therapy (VMAT) plans for total scalp irradiation. We selected a single adult model case and compared the dosimetric results for the three plans. All plans mainly used 6-MV photon beams, and the prescription dose was 60 Gy in 30 fractions. First, we compared the LPE, HT and VMAT plans, with all plans including a 1-cm bolus. We also compared HT plans with and without the bolus. The conformity indices for LPE, HT and VMAT were 1.73, 1.35 and 1.49, respectively. The HT plan showed the best conformity and the LPE plan showed the worst. However, the plans had similar homogeneity indexes. The dose to the hippocampus was the highest in the VMAT plan, with a mean of 6.7 Gy, compared with 3.5 Gy in the LPE plan and 4.8 Gy in the HT plan. The doses to the optical structures were all within the clinically acceptable range. The beam-on time and monitor units were highest in the HT plan. The HT plans with and without a bolus showed similar target coverage and organ-at-risk (OAR) sparing. The HT plan showed the best target coverage and conformity, with low doses to the brain and hippocampus. This plan also had the advantage of not necessarily requiring a bolus. Although the VMAT plan showed better conformity than the LPE plan and acceptable OAR sparing, the dose to the hippocampus should be considered when high doses are prescribed.

KEYWORDS: total scalp irradiation; lateral photon–electron technique; helical tomotherapy; volumetric-modulated arc therapy, hippocampus

INTRODUCTION

Total scalp irradiation is used to treat several conditions such as lymphoma, Kaposi’s sarcoma, angiosarcoma, and extensive skin cancer [1–5]. Historically, homogenous scalp irradiation has presented technical and dosimetric challenges because of the complex shape and superficial nature of the target.

Several techniques have been suggested for conventional linear-accelerator treatment. Because of the relatively high surface dose provided by electrons, techniques using several electron beams have been developed [1, 6, 7]. However, several problems have been associated with the use of multiple electron beams, including dose heterogeneity across the target, numerous field junctions, and...
laborious treatment setup. To overcome these disadvantages, Akazawa incorporated a megavoltage photon beam and introduced the lateral photon–electron technique (LPE) [8]. This technique incorporates parallel opposing lateral photon beams, which cover the outer rind of the scalp around the top of the head, and lateral electron fields, which cover the remaining lateral portion of the scalp, matched with the photon beam. Tung et al. overlapped the photon and electron fields to improve dose uniformity [9]. Kinard et al. described a photon shell technique with four coplanar arcs [10].

With the recent advancement of radiation delivery machines, several newer modalities have been used for total scalp irradiation. Intensity-modulated radiation therapy (IMRT), high-dose-rate (HDR) brachytherapy, and serial tomotherapy have been studied by several investigators [2, 11–13]. Wojcicka et al. compared step-and-shoot IMRT and HDR brachytherapy with LPE, concluding that IMRT provided the best coverage and homogeneity. HDR was the most conformal plan, but the total dose delivered was limited because of the brain and optical structures [13]. Locke et al. compared serial tomotherapy with LPE, showing that LPE was superior because serial tomotherapy was associated with a substantial dose to the brain and optical structures [11].

Helical tomotherapy (HT) is a form of computed tomography (CT)-guided IMRT that uses 6-MV beams modulated by a 64-leaf binary multileaf collimator (MLC) [12]. HT is well suited to scalp irradiation because of its ability to deliver a tangential beam to any point on the scalp. It also does not have a field-matching problem. Another newer form of IMRT based on volumetric-modulated arc therapy (VMAT) has also become clinically available. VMAT machines can deliver radiation consisting of one or more arcs using a varying dose rate, changing gantry rotation speed, and bidirectional MLC motion [14]. VMAT also appears to be a good modality for scalp irradiation because it shares some of the advantages of HT, yet also offers a shorter treatment time. This reduced treatment time is possible because VMAT involves a cone beam, whereas HT involves a fan beam.

In this study, we selected a single adult model case for total scalp irradiation and performed a treatment-planning study to compare the HT and VMAT plans with the LPE plan, the most commonly used technique in conventional plans for total scalp irradiation. Several dosimetric parameters were compared, and we additionally focused on hippocampus sparing, which has recently been recognized as an important issue for neurocognitive function when treating the brain area [15].

MATERIALS AND METHODS
IRB approval
We received approval for this study from the Institutional Review Board of the Incheon St. Mary’s Hospital, the Catholic University of Korea (Reference No. OC13RIS0016).

Target delineation and prescription
We selected an adult patient without a visible gross tumor on imaging studies. A CT scan was obtained while the patient was immobilized using a thermoplastic head mask. The CT images were acquired in 2.5-mm thick slices. The target volume and normal structures were delineated by a single experienced physician.

The target was delineated on the whole scalp from the surface of the scalp skin to the depth of the cranium. The organs at risk (OARs) used for optimization included the brain, brain stem, hippocampus, eyeballs, lens, and optic nerves. The hippocampus was contoured based on the fusion of magnetic resonance imaging (MRI) scans with simulation CT scans and the contouring guideline suggested by Gondi et al. [16]. A hippocampal avoidance region was also created using a 5-mm volumetric expansion around the hippocampus, and this OAR was used for the planning and dosimetric analyses.

The prescription dose was 60 Gy to the target in 30 fractions. The plans were optimized to achieve a target coverage with at least 90% of the planning target volume (PTV) receiving 100% of the prescription dose ($V_{60Gy} > 90\%$) and to minimize the dose to the OARs. To achieve these planning goals, a 1-cm thick bolus was needed for the LPE and VMAT plans. It was impossible to achieve a $V_{60Gy} > 90\%$ without a bolus for the LPE and VMAT plans. However, a bolus was not essential for the HT plan to achieve the planning goal. This study employed the use of a virtual bolus created by the planning system (Eclipse ver. 8.9) rather than an actual bolus.

The LPE, HT and VMAT treatment plans
The LPE plan consisted of opposing lateral photon and electron beams, as previously described by Akazawa [8]. The outer field comprised 6-MV photon beams with a field size of $22 \times 14\text{ cm}^2$. The inner field comprised 9-MeV electron beams with a cone size of 25 cm. The electron beam was matched to the photon field on the skin (Fig. 1a). For the LPE plan, a 1-cm thick virtual bolus was applied. A pencil-beam algorithm was used for the photon field calculations, and an electron Monte Carlo algorithm was used for the electron field, as provided by the Eclipse ver. 8.9 treatment-planning system. The calculation grid was $2.5 \times 2.5 \text{ mm}$. The Tomotherapy Hi-Art system (Tomotherapy Inc., Madison, WI, USA) is an IMRT modality in which a 6-MV linear accelerator is mounted on a ring gantry that continuously rotates during treatment. The couch simultaneously moves through the rotating beam plane so that the radiation is delivered in a helical fashion [17]. For HT planning, three parameters should be selected: the field width (the slice thickness of the radiation field projected at the isocenter along the gantry rotation axis) was set to 1.0 cm, the pitch (the couch movement relative to the field width during one gantry rotation) was set to 0.172, and the modulation factor (the ratio between the maximum number of opening leaves and the average number of opening leaves in active gantry rotation) was set to 3.20. The bolus was not essential for HT planning. Accordingly, we generated two plans: a plan with the bolus and a plan without the bolus. Tomotherapy Planning Station ver. 4.0.4 was used, and the calculation grid was $1.95 \times 1.95 \text{ mm}$.

RapidArc (Varian Medical Systems, Palo Alto, CA, USA) treatment was planned with the Eclipse ver. 10.0.42 planning system. The plans were optimized using the Progressive Resolution Optimizer (PRO) ver. 10.0.28 and calculated with the Anisotropic Analytical Algorithm (AAA) ver. 10.0.28. It was impossible to achieve proper target coverage with single- or double-isocenter plans. Therefore, the final approved plan consisted of three centers that were located 4 cm apart. The locations of the three centers are shown in Fig. 1b. A total of nine 6-MV clockwise 358° arcs (181° to 179°) were used with no treatment table rotation. The collimator angles were set to 90° for each arc, and the MLC margin was 0.5 cm from the target.
VMAT plan, a 1-cm thick bolus was used. The calculation grid was 2.5 × 2.5 mm.

Analysis and comparisons between plans
Plan conformity and homogeneity were assessed using the following parameters.

Conformity index
The conformity index (CI) is a ratio used to evaluate how well the target fits the prescription isodose volume in the plan [18]:

\[ CI = \frac{V_{PTV}}{TV_{PTV}} \times \frac{V_{TV}}{TV_{TV}}; \]

where \( V_{PTV} = \) PTV volume, \( V_{TV} = \) volume of the prescribed isodose lines, \( TV_{PTV} = \) volume of \( V_{PTV} \) within the \( V_{TV} \). A lower CI value indicates better conformity.

Homogeneity index
The homogeneity index (HI) is a ratio used to evaluate dose homogeneity in the PTV [19, 20]:

\[ HI = \frac{D_{1\%}}{D_{99\%}}; \]

where D1% and D99% are the minimum doses delivered to 1% and 99% of the PTV. A lower HI value indicates better homogeneity.

Conformation number
The conformation number (CN) is an index that incorporates both target coverage and the extent to which the normal tissues are spared. CN was introduced by vant’ Riet et al [21].

\[ CN = \frac{TV_{R1}}{TV} \times \frac{TV_{R1}}{V_{R1}}; \]

where \( TV = \) target volume, \( TV_{R1} = \) target volume covered by the reference dose, and \( V_{R1} = \) volume of the reference dose.

Conformation index
The conformation index (COIN) is an index that was proposed by Baltas et al. and is computed for a reference dose [22]:

\[ COIN = CN \times \prod_{i=1}^{NCO} \left[ 1 - \frac{V_{COI,i}}{V_{CO,i}} \right], \]

where \( CN = \) conformation number, \( NCO = \) number of critical organs, \( V_{COI,i} = \) critical volume receiving at least the reference dose, and \( V_{CO,i} = \) critical organ volume.

‘Index A’ improvement ratio
The ‘Index A’ improvement ratio (\( \Delta Index A (\%) \)) is a ratio used to evaluate the improvement of ‘Index A’ between the plans. It was employed for the CI, HI, CN and COIN indices [19]:

\[ \Delta Index A (\%) = \frac{Index A_{plan2} - Index A_{plan1}}{Index A_{plan1}} \times 100\%. \]

Quality index
The quality index (QI) is an index used to evaluate differences in the absorbed doses at the OARs. This index uses the maximum dose for serial OARs (optic nerves in our study) and the mean dose for parallel OARs (all other OARs in our study) [20].

\[ QI_{Serial} = \frac{D_{plan1}}{D_{max1}}; \quad QI_{Parallel} = \frac{D_{plan1}}{D_{mean1}}. \]

We compared the LPE, HT and VMAT plans by assessing these parameters, which were all planned with a 1-cm thick bolus. The beam-on time and monitor units (MUs) of each system were also compared. Because an optimal HT plan could be obtained without the use of a bolus, we also compared HT plans with and without the bolus.
RESULTS
Comparisons of the LPE, HT and VMAT plans with a bolus

Isodose curves and the dose–volume histograms (DVHs) of the target are shown in Figs 2 and 3. Table 1 lists the indices that indicate the conformity and homogeneity of the target coverage in each plan. The CI was 1.73 with LPE, 1.35 with HT, and 1.49 with VMAT, indicating that the HT plan had the best conformity and the LPE plan had the worst. The CI was improved by 22.0% in the HT plan and 13.9% in the VMAT plan compared with the LPE plan. The HT plan showed 10.4% better CI compared with the VMAT plan. While the CI clearly differed between the plans, the HI was similar across plans. Homogeneity was poorest for the VMAT plan; however, the difference was only 1.7%.

The CN and COIN results resembled the CI results. The CN and COIN were highest in the HT plan, second highest in the VMAT plan and lowest in the LPE plan. These findings suggest that the HT plan had the best conformity and the smallest volume of normal structures included in the high-dose area. The difference between the LPE and HT plans was > 25%, that between the LPE and VMAT plans was ~17%, and that between the HT and VMAT plans was 10.8%.

Dosimetric comparisons of the OARs are shown in Tables 2 and 3. The LPE plan featured the highest mean and maximum dose to the brain, as well as the largest brain volumes that received at least 20 Gy, 30 Gy, 40 Gy and 50 Gy (V20Gy, V30Gy, V40Gy and V50Gy). However, as shown in Fig. 3, the DVHs for LPE and VMAT crossed at the 15-Gy level, indicating that the volume receiving < 15 Gy was larger in the VMAT plan.

Other normal structures, including the hippocampus, received the lowest mean and maximum doses in the LPE plan (Table 2, Fig. 3). However, all doses to normal structures were within clinically acceptable levels in the HT and VMAT plans. The mean dose to the hippocampus was 3.5 Gy in the LPE plan, 4.8 Gy in the HT plan and 6.7 Gy in the VMAT plan.

Comparing the CI values of the LPE plan with the others, the CI values of all the OARs except for the brain were over 1.0. This result indicates that all OARs except the brain received a lower dose in the LPE plan than they did in the other plans. The mean CI values were 1.78 ± 0.74 for the HT versus LPE plan, and 2.73 ± 0.90 for the VMAT versus LPE plan. However, the respective brain CI values were 0.58 and 0.86.

The CI values showed that all nine OARs received lower doses in the HT plan compared with the VMAT plan. The mean CI value for VMAT versus HT was 1.60 ± 0.38.

The beam-on times and MUs for a single fraction (to deliver 2 Gy to the PTV) are listed in Table 4. The LPE plan had the shortest beam-on time. The beam-on times were 1.5 min for the LPE plan,
14.7 min for the HT plan and 11.3 min for the VMAT plan. The MUs were also the lowest in the LPE plan (616). The MUs for the HT plan were 13,088, which was much higher than the MUs of 1,791 in the VMAT plan. HT and VMAT differed much more substantially in terms of MUs than they did in terms of beam-on time.

Comparisons of HT plans with and without a bolus

For LPE and VMAT, it was impossible to obtain acceptable plans that satisfied the planning goal without using a bolus. However, we were able to obtain a well-designed HT plan without a bolus. Tables 5 and 6 present the target and OAR analyses for the HT plans with and without a bolus.

### Table 1. Dosimetric comparisons of target coverage in the LPE, HT and VMAT plans

|        | LPE | HT  | VMAT | HT/LPE | VMAT/LPE | VMAT/HT |
|--------|-----|-----|------|--------|----------|---------|
| CI     | 1.73| 1.35| 1.49 | -22.0  | -13.9    | 10.4    |
| HI     | 1.15| 1.15| 1.17 | 0      | 1.7      | 1.7     |
| CN     | 0.57| 0.74| 0.66 | 29.8   | 15.8     | -10.8   |
| COIN   | 0.56| 0.74| 0.66 | 32.1   | 17.9     | -10.8   |

LPE = lateral photon–electron plan, HT = helical tomotherapy, VMAT = volumetric-modulated arc therapy, CI = conformity index, HI = homogeneity index, CN = conformation number, COIN = conformation index.
without a bolus. The CI, HI, CN and COIN were similar between the plans. The CI, CN and COIN were 6.7% better in the HT plan without the bolus, suggesting better conformity. The HI was also 0.8% better in the HT plan without the bolus.

With respect to OAR sparing, the mean QI value was 1.08 ± 0.14, which suggested slightly better OAR sparing in the HT plan with the bolus. While the brain stem, left eyeball and hippocampus received a slightly lower dose in the plan without the bolus, the mean dose differences were only 0.3 Gy in the brain stem and 0.1 Gy in the hippocampus. Although the doses to normal structures were all in the clinically acceptable range in both plans, the DVHs showed that the brain volume was lower for all dose ranges (V10Gy, V20Gy, V30Gy, V40Gy and V50Gy) in the plan with the bolus (Fig. 4). However, this only amounted to a 0.5 Gy difference from the mean brain dose.

**DISCUSSION**

Total scalp irradiation is relatively rare in clinical practice. However, there are several diseases for which total scalp irradiation is indicated, including lymphoma, mycosis fungoides, Kaposi’s sarcoma, angiosarcoma, and extensive skin cancer [1–5]. The prescription...
dose is variable, depending on the disease category and the purpose of radiation treatment. For example, 20–30 Gy is usually prescribed for mycosis fungoides and other lymphomas, 25–40 Gy for Kaposi's sarcoma, and 50–60 Gy for skin cancer and angiosarcoma. This dose variability and the complex shape of the scalp make it difficult to select a single best technique for all patients. In this study, a detailed dosimetric analysis was performed to compare three different techniques (the conventional LPE plan, an HT plan, and a VMAT plan) for an assumed prescription dose of 60 Gy to the scalp.

The most commonly used technique is LPE, which was first described by Akazawa [8]. From our results, the advantage of this technique over the other two techniques was that the optical structures received the lowest dose. The mean doses to several optical structures were all below 2 Gy. However, the mean brain dose was higher for the HT and VMAT plans. In addition to the highest mean brain dose, LPE was also associated with the highest $V_{20Gy}$, $V_{30Gy}$, $V_{40Gy}$, and $V_{50Gy}$. With respect to target coverage, the LPE plan was the least conformal, although the homogeneity was similar across plans. The technical difficulty of matching the photon-electron beams was found to be another disadvantage of LPE.

The VMAT plan showed better dose conformity than the LPE plan; the conformity increased by 13.9% compared with LPE. However, the doses to the OARs were the highest in this plan, with the exception of the brain dose. The mean QI values were 2.73 compared with LPE, and 1.6 compared with HT. Nevertheless, the doses to the optical structures were within the clinically acceptable range. The mean brain dose was higher than that of HT, but lower than that of LPE.

The HT plan showed the best conformity among the plans. The conformity was increased by 10.4% compared with VMAT and 22.0% compared with LPE. The mean dose to the brain was also the lowest. Other optical structure doses were higher than those of LPE, but lower than those of VMAT. This superiority of HT compared with LPE was consistent with other studies. Orton et al. compared the HT plan with two conventional linear accelerator plans (LPE and concentric electron fields technique) [12]. They prescribed 40 Gy in 20 fractions, and found that the HT plan delivered a more uniform dose to the scalp and reduced the $V_{30Gy}$ of the brain by as much as two-thirds.

Table 5. Dosimetric comparisons of target coverage for HT plans with and without a bolus

| OARs      | HT with a bolus | HT without a bolus | HT without a bolus/HT with a bolus |
|-----------|----------------|-------------------|-----------------------------------|
| CI        | 1.35           | 1.26              | ΔCI (%) −6.7                       |
| HI        | 1.15           | 1.14              | ΔHI (%) −0.8                       |
| CN        | 0.74           | 0.79              | ΔCN (%) 6.7                        |
| COIN      | 0.74           | 0.79              | ΔCOIN (%) 6.7                      |

HT = helical tomotherapy, CI = conformity index, HI = homogeneity index, CN = conformation number, COIN = conformation index.

Table 6. Dosimetric comparisons of the mean, maximum and minimum doses to nine OARs for HT plans with and without a bolus

| OARs     | Mean (Gy) | Max (Gy) | Min (Gy) | QI value |
|----------|-----------|----------|----------|----------|
|          | With a bolus | Without a bolus | With a bolus | Without a bolus | With a bolus | Without a bolus |
| Brain    | 9.8       | 10.3     | 57.4     | 59.5     | 1.0        | 0.9        | 1.05        |
| Brain stem | 3.6      | 3.3      | 5.7      | 4.4      | 1.5        | 1.4        | 0.92        |
| Hippocampus | 4.8    | 4.7      | 6.5      | 6.4      | 4.7        | 3.5        | 0.98        |
| Lt eyeball | 5.2     | 4.8      | 18.7     | 20.9     | 1.8        | 2.1        | 0.92        |
| Rt eyeball | 4.5    | 6.0      | 17.3     | 19.7     | 1.7        | 1.7        | 1.33        |
| Lt lens   | 3.0       | 3.7      | 3.6      | 4.5      | 2.4        | 3.1        | 1.23        |
| Rt lens   | 2.4       | 2.7      | 3.0      | 3.4      | 2.0        | 2.2        | 1.12        |
| Lt optic nerve | 3.5  | 3.8      | 5.0      | 5.8      | 2.8        | 3.3        | 1.08        |
| Rt optic nerve | 2.9 | 3.2      | 3.6      | 3.9      | 2.4        | 2.7        | 1.10        |
| $V_{10Gy}$ (%) |         |          |          |          | Brain     | 25.0       | 25.4       | 12.5       | 13.9       | 7.4        | 8.8        |
| $V_{20Gy}$ (%) |         |          |          |          | Brain     | 3.0        | 4.4        | 0.3        | 1.1        |

OARs = organs at risk, QI = quality index, Lt = left side, Rt = right side, $V_{10Gy} = \text{volume receiving at least } 10 \text{ Gy}, V_{20Gy} = \text{volume receiving at least } 20 \text{ Gy}, V_{30Gy} = \text{volume receiving at least } 30 \text{ Gy}, V_{40Gy} = \text{volume receiving at least } 40 \text{ Gy}, V_{50Gy} = \text{volume receiving at least } 50 \text{ Gy}$. 

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Another advantage of the HT plan was that the bolus was not an essential component for planning. Unlike the LPE and VMAT plans, in which it was impossible to achieve appropriate target coverage without a bolus, the HT plans with and without the bolus did not differ markedly. It is also practically difficult to apply the bolus tightly around the entire scalp. The advantage of the bolus in the HT plan was only a slight decrease in the brain dose. The mean dose decreased by only 0.5 Gy (from 10.3 to 9.8 Gy) when 60 Gy was prescribed to the scalp. Several studies have shown prescribed doses can be delivered with HT, even to regions close to the surface, without the use of a bolus [17, 23–25]. Although the results were variable across these studies, the measured surface doses were not that different from the calculated doses, and this can be explained by the high number of tangential beams, which is unique to HT [17].

On the other hand, for cases in which the skin surface does not need to receive therapeutic doses, the VMAT plan can also be a good option. Kelly et al. compared static IMRT with VMAT for total dural irradiation [26]. They cropped 3 mm of the PTV to allow skin sparing, and concluded that VMAT can provide a clinically acceptable treatment plan without the use of a bolus.

However, the HT plan had the longest beam-on time and the highest MUs. The HT beam-on time was 9.8 and 1.3 times longer than the beam-on times of the LPE and VMAT plans, respectively. Although the HT plan showed only a slightly longer beam-on time than the VMAT plan, the MUs were 21.2 and 7.3 times higher than the MUs of the LPE and VMAT plans, respectively. Treatments with longer delivery times generally have the potential disadvantage of intrafraction patient motion, and higher total MU delivery may result in higher patient exposure to leakage radiation [27]. This concern regarding HT is common for other treatment sites [27, 28].

Clinically, the main concern while treating the scalp is the dose to the brain. The maximal dose to the brain was higher than 60 Gy only...
in the LPE plan. The brain dose was not high enough to cause brain necrosis in the HT or VMAT plans. However, neurocognitive function could be affected, even at low doses [29]. Recently, several clinical studies have suggested that radiation damage to the hippocampus plays a considerable role in neurocognitive function decline [15]. Deficits in learning, memory, and spatial processing are thought to be related to hippocampus injury in patients who have received whole brain radiation. However, a widely accepted dose guideline does not yet exist. Gondi et al. suggests that irradiation of 40% of the bilateral hippocampus at a dose of > 7.3 Gy is associated with long-term impairment after benign and low-grade adult brain tumor treatment [30]. Hsu et al. and Gutierrez et al. studied the feasibility of VMAT and HT for the hippocampal sparing whole brain radiotherapy with a simultaneous integrated boost. The dose constraint used for the hippocampus was 6 Gy in both studies [31, 32]. In another study, Gondi et al. suggests that, when whole brain radiotherapy is delivered with 30 Gy in 10 fractions, it is able to reduce (i) the mean dose to the hippocampus by 5.5 Gy using HT and by 7.8 Gy using LINAC-based IMRT, and (ii) the maximum dose to 12.8 Gy using HT and to 15.3 Gy using LINAC-based IMRT [15]. In the Radiation Therapy Oncology Group (RTOG) 0933 Phase II trial for hippocampus sparing in brain metastasis treatment, the guidelines are as follows: (i) dose to 100% of the hippocampal volume ($D_{100}$) $\leq 10$ Gy and (ii) maximum dose $\leq 17$ Gy. In our study, the VMAT plan showed the highest dose to the hippocampus, with a mean dose of 6.7 Gy (as compared with 3.5 Gy for LPE and 4.8 Gy for HT). The maximum dose was 9.8 Gy in the VMAT plan, which was higher than the 4.6 Gy dose for LPE and the 6.5 Gy dose for HT. The doses were smaller than those noted in previous studies. However, considering the hypothesis that active neural stem cells in the hippocampal dentate gyrus play a major role in neurocognitive function and that these cells are quite sensitive to radiation [29], lowering the dose to the hippocampus would be needed.

In conclusion, the results of our study showed that HT was the best modality for total scalp irradiation. The HT plan showed the best target coverage and conformity, with low doses to the brain and hippocampus. It also has the advantage of not requiring a bolus. Although the VMAT plan showed acceptable OAR sparing and better conformity compared with the LPE plan, the dose to the hippocampus should be considered when high doses ($> 60$ Gy) are needed. The LPE plan has the advantage of lowering the dose to the optical structures. However, considering the LPE’s conformity limitations, the large volume of the brain that received high doses, and the technical difficulty associated with matching the fields, it seems as though LPE should cede its throne to other more recent modalities.

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REFERENCES
1. Mollenberg DE, Schoeppel SL. Total scalp treatment of mycosis fungoides: the 4 x 4 technique. Int J Radiat Oncol Biol Phys 1993;27:953–8.
2. Bedford JL, Childs PJ, Hansen VN, et al. Treatment of extensive scalp lesions with segmental intensity-modulated photon therapy. Int J Radiat Oncol Biol Phys 2005;62:1549–58.
3. Samant RS, Fox GW, Gerig LH, et al. Total scalp radiation using image-guided IMRT for progressive cutaneous T cell lymphoma. Br J Radiol 2009;82:122–5.
4. Guadagnolo BA, Zagars G, Araujo D, et al. Outcomes after definitive treatment for cutaneous angiosarcoma of the face and scalp. Head Neck 2011;33:661–7.
5. Stelzer KJ, Griffin TW. A randomized prospective trial of radiation therapy for AIDS-associated Kaposi’s sarcoma. Int J Radiat Oncol Biol Phys 1993;27:1057–61.
6. Able CM, Mills MD, McNeese MD, et al. Evaluation of a total scalp electron irradiation technique. Int J Radiat Oncol Biol Phys 1991;21:1063–72.
7. Walker C, Wadd NJ, Lucaffa HH. Novel solutions to the problems encountered in electron irradiation to the surface of the head. Br J Radiol 1999;72:787–91.
8. Akazawa C. Treatment of the scalp using photon and electron beams. Med Dosim 1989;14:129–31.
9. Tung SS, Shiu AS, Starkschall G, et al. Dosimetric evaluation of total scalp irradiation using a lateral electron-photon technique. Int J Radiat Oncol Biol Phys 1993;27:153–60.
10. Kinard JZ, Zwicker RD, Schmidt Ullrich RK, et al. Short communication: Total craniofacial photon shell technique for radiotherapy of extensive angiosarcomas of the head. Br J Radiol 1996;69:351–5.
11. Locke J, Low DA, Grigireit T, et al. Potential of tomotherapy for total scalp treatment. Int J Radiat Oncol Biol Phys 2002;52:553–9.
12. Orton N, Jaradat H, Welsh J, et al. Total scalp irradiation using helical tomotherapy. Med Dosim 2008;33:162–8.
13. Wojcicka JB, Lasher DE, McAfee SS, et al. Dosimetric comparison of three different treatment techniques in extensive scalp lesion irradiation. Radiother Oncol 2009;91:255–60.
14. Otto K. Volumetric modulated arc therapy: IMRT in a single gantry arc. Med Phys 2008;35:310–7.
15. Gondi V, Tome WA, Mehta MP. Why avoid the hippocampus? A comprehensive review. Radiother Oncol 2010;97:370–6.
16. Gondi V, Tolakanahalli R, Mehta M, et al. Hippocampal-sparing whole-brain radiotherapy: a “how-to” technique using helical tomotherapy and linear accelerator-based intensity-modulated radiotherapy. Int J Radiat Oncol Biol Phys 2010;78:1244–52.
17. Avanço M, Drigo A, Ren Kaiser S, et al. Dose to the skin in helical tomotherapy: results of in vivo measurements with radiochromic films. Phys Med 2013;29:304–11.
18. Nakamura JL, Verhey LJ, Smith V, et al. Dose conformity of gamma knife radiosurgery and risk factors for complications. Int J Radiat Oncol Biol Phys 2001;51:1313–9.
19. Lee TF, Chao PJ, Wang CY, et al. Dosimetric comparison of helical tomotherapy and dynamic conformal arc therapy in stereotactic radiosurgery for vestibular schwannomas. Med Dosim 2011;36:62–70.
20. Sheng K, Molloy JA, Larner JM, et al. A dosimetric comparison of non-coplanar IMRT versus Helical Tomotherapy for nasal cavity and paranasal sinus cancer. Radiother Oncol 2007;82:174–8.
21. van’t Riet A, Mak AC, Moerland MA, et al. A conformation number to quantify the degree of conformity in brachytherapy
and external beam irradiation: application to the prostate. *Int J Radiat Oncol Biol Phys* 1997;37:731–6.

22. Baltas D, Kolotas C, Geramani K, et al. A conformal index (COIN) to evaluate implant quality and dose specification in brachytherapy. *Int J Radiat Oncol Biol Phys* 1998;40:515–24.

23. Ramsey C, Seibert R, Robison B, et al. Helical tomotherapy superficial dose measurements. *Med Phys* 2007;34:3286–93.

24. Tournel K, Verellen D, Duchateau M, et al. An assessment of the use of skin flashes in helical tomotherapy using phantom and in-vivo dosimetry. *Radiother Oncol* 2007;84:574–81.

25. Zibold F, Sterzing F, Sroka Perez G, et al. Surface dose in the treatment of breast cancer with helical tomotherapy. *Strahlenther Onkol* 2009;185:574–81.

26. Kelly PJ, Mannarino E, Lewis JH, et al. Total dural irradiation: RapidArc versus static-field IMRT: a case study. *Med Dosim* 2012;37:175–81.

27. Caudell JJ, De Los Santos JF, Keene KS, et al. A dosimetric comparison of electronic compensation, conventional intensity modulated radiotherapy, and tomotherapy in patients with early-stage carcinoma of the left breast. *Int J Radiat Oncol Biol Phys* 2007;68:1505–11.

28. Van Gestel D, van Vliet-Vroegindeweij C, Van den Heuvel F, et al. RapidArc, SmartArc and TomoHD compared with classical step and shoot and sliding window intensity modulated radiotherapy in an oropharyngeal cancer treatment plan comparison. *Radiother Oncol* 2013;8:37.

29. Mizumatsu S, Monje M, Morhardt D, et al. Extreme sensitivity of adult neurogenesis to low doses of X-irradiation. *Cancer Res* 2003;63:4021–7.

30. Gondi V, Hermann BP, Mehta MP, et al. Hippocampal dosimetry predicts neurocognitive function impairment after fractionated stereotactic radiotherapy for benign or low-grade adult brain tumors. *Int J Radiat Oncol Biol Phys* 2013;85:348–54.

31. Gutierrez AN, Westerly DC, Tome WA, et al. Whole brain radiotherapy with hippocampal avoidance and simultaneously integrated brain metastases boost: a planning study. *Int J Radiat Oncol Biol Phys* 2007;69:589–97.

32. Hsu F, Carolan H, Nichol A, et al. Whole brain radiotherapy with hippocampal avoidance and simultaneous integrated boost for 1–3 brain metastases: a feasibility study using volumetric modulated arc therapy. *Int J Radiat Oncol Biol Phys* 2010;76:1480–5.