Case Study

Linac-based stereotactic radiosurgery for the treatment of multiple brain metastases

Institution
Geetanjali Cancer Center

Location
Geetanjali University, Udaipur, Rajasthan, India

Medical Staff
Dr Shankar Vangipuram
Director, Division of Radiosurgery

Mr Vinoth Kumar
Chief Medical Physicist
A 55 year old female presented in May 2015 with symptoms of increased intracranial pressure (ICP), including headache, nausea and vomiting. The patient was previously diagnosed with breast cancer in 2013. In February 2014 she had a radical mastectomy. Subsequently she received chemotherapy (four cycles of AC and four cycles of Taxol) followed by radiotherapy to the chest wall.
Since this patient had triple negative breast cancer (negative for estrogen receptors, progesterone receptors and HER2/neu), the presence of brain metastases was suspected. An MRI scan of the patient’s brain revealed 3 brain metastases located in the right cerebellum, the left temporal lobe and the high frontal lobe.

The patient had no other comorbidities, making her a good candidate for stereotactic radiosurgery (SRS) according to the standard protocol for brain metastases at this institution (Figure 1), which is based on ASTRO recommendations1,2, IRSA guidelines3 and recent published evidence 4-6.

Summary

Patient demographics
55 year old woman treated in 2013 for breast cancer and presenting with symptoms of increased intracranial pressure (ICP).

Treatment
Stereotactic radiosurgery (SRS). Marginal dose prescribed per lesion: GTV1 18 Gy; GTV2 20 Gy; GTV3 14 Gy, delivered in a single fraction using a common isocenter and a double VMAT arc.

Diagnosis
3 brain metastases detected by MRI, located in the right cerebellum, the left temporal lobe and the right frontal lobe with dimensions: GTV1 0.82 x 1.01 x 1.15 cm; GTV2 1.07 x 1.62 x 1.42 cm; GTV3 2.58 x 2.48 x 3.8 cm.

Treatment Planning and Delivery System
- Monaco® treatment planning system version 5.0.1.
- Versa HD™ with Agility™ MLC
- Fraxion™ cranial immobilization
- HexaPOD™ evo RT robotic couch
- XVI image guidance
There is a strong case for performing SRS as a primary treatment for patients with 2–4 metastases and good functional status in order to reduce the risk of neurocognitive decline associated with whole brain radiotherapy (WBRT). The treatment modality used for SRS should be fast, to reduce the time that the patient must remain still on the couch, and its target-hitting precision must be very robust, to spare healthy tissue and critical structures in the brain. Such speed and precision can now be achieved with the Versa HD linear accelerator with Agility multileaf collimator (MLC), using the Monaco treatment planning system (TPS).

The combination of Monaco, with its advanced dose modulation capabilities and its ability to achieve excellent dose distributions, and Agility, with its rapid leaf speed, low transmission and high-resolution beam shaping, allows very accurate dose placement. This is particularly important for the high doses per fraction required for stereotactic approaches and can be achieved, even for small targets and complex plans, using these tools. In addition, the large field size and interdigitation capabilities of Agility allow plans to be generated for multiple targets using a single isocenter, which has the desirable effect of reducing treatment time and Monitor Units to the patient.
Treatment planning

For treatment planning and treatment delivery the patient was immobilised using the Elekta Fraxion cranial immobilization system with a thermoplastic mask. CT and MRI scans (T1 post contrast, T2 and proton density sequences) were obtained, each with 1.25 mm slice thickness, and fused prior to image registration and delineation of target volumes and critical structures (Figure 2).

A margin of 2 mm was applied to the gross tumour volumes GTV1, GTV2 and GTV3 in all directions, except for the superior and inferior sides, to form planning target volumes PTV1, PTV2 and PTV3 respectively. The volume, dimensions and dose prescription for each volume is shown in Table 1. Dose constraints for critical structures are shown in Table 2 and the dose volume histogram (DVH) for target volumes and critical structures are shown in Figure 3.
### Table 1.
Dimensions and prescribed dose for each of the 3 brain metastases

<table>
<thead>
<tr>
<th>Volume name</th>
<th>Volume (cc)</th>
<th>Dimensions (cm)</th>
<th>Dose Prescription</th>
</tr>
</thead>
<tbody>
<tr>
<td>GTV1</td>
<td>0.446</td>
<td>0.82 x 1.01 x 1.15</td>
<td>18 Gy</td>
</tr>
<tr>
<td>PTV1</td>
<td>0.799</td>
<td>1.05 x 1.01 x 1.15</td>
<td></td>
</tr>
<tr>
<td>GTV2</td>
<td>0.953</td>
<td>1.07 x 1.62 x 1.42</td>
<td>20 Gy</td>
</tr>
<tr>
<td>PTV2</td>
<td>1.461</td>
<td>1.37 x 1.62 x 1.42</td>
<td></td>
</tr>
<tr>
<td>GTV3</td>
<td>15.144</td>
<td>2.58 x 2.48 x 3.8</td>
<td>14 Gy</td>
</tr>
<tr>
<td>PTV3</td>
<td>18.252</td>
<td>2.8 x 2.48 x 4.8</td>
<td></td>
</tr>
</tbody>
</table>

### Table 2.
Dimensions and prescribed dose for critical structures

<table>
<thead>
<tr>
<th>Critical structure</th>
<th>Volume (cm³)</th>
<th>Min dose (cGy)</th>
<th>Max dose (cGy)</th>
<th>Mean dose (cGy)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brain stem</td>
<td>26.645</td>
<td>34.7</td>
<td>827.6</td>
<td>341.2</td>
</tr>
<tr>
<td>Left eye</td>
<td>10.444</td>
<td>15.3</td>
<td>323.6</td>
<td>146.2</td>
</tr>
<tr>
<td>Right eye</td>
<td>9.550</td>
<td>9.4</td>
<td>318.0</td>
<td>174.4</td>
</tr>
<tr>
<td>Left hippocampus</td>
<td>2.255</td>
<td>127.5</td>
<td>593.1</td>
<td>348.1</td>
</tr>
<tr>
<td>Right hippocampus</td>
<td>2.328</td>
<td>225.5</td>
<td>523.5</td>
<td>357.8</td>
</tr>
<tr>
<td>Left optical nerve</td>
<td>0.563</td>
<td>118.8</td>
<td>369.9</td>
<td>244.4</td>
</tr>
<tr>
<td>Right optical nerve</td>
<td>0.573</td>
<td>188.6</td>
<td>395.6</td>
<td>255.0</td>
</tr>
<tr>
<td>Optic chiasma</td>
<td>2.469</td>
<td>212.9</td>
<td>619.1</td>
<td>403.4</td>
</tr>
<tr>
<td>Pituitary</td>
<td>0.972</td>
<td>287.5</td>
<td>629.3</td>
<td>483.8</td>
</tr>
<tr>
<td>Patient (unspecified tissue)</td>
<td>133.820</td>
<td>0.2</td>
<td>845.8</td>
<td>50.4</td>
</tr>
</tbody>
</table>
A VMAT plan was generated using a single, common isocenter for all three targets. The isocenter was placed at the center of the volume containing all three target volumes. The distances between the isocenter and the plane containing the central slice of each PTV are 2.62 cm for PTV1, 0.7 cm for PTV2 and 1.70 cm for PTV3. An optimal plan was generated using a 6 MV flattened beam using Monaco version 5.0.1 and Agility, which was reviewed by the Head of Radiation Oncology prior to treatment delivery. Dose coverage for each target volume was comparable to an equivalent plan generated using three separate isocenters (Figure 3/Figure 4).

Figure 3. Dose Volume Histogram (DVH) for target volumes and critical structures using a single isocenter and multiple isocenters.
Figure 4.
Comparison of single and multiple isocenter plans
In addition, excellent dose fall-off was achieved as demonstrated by measuring the distance between the 100% and 80% isolines and between the 100% and 50% isolines (Table 4). This is particularly important for normal brain tissue and critical structure sparing in SRS.

<table>
<thead>
<tr>
<th>Target volume</th>
<th>Max distance between 100% and 80% isoline (mm)</th>
<th>Max distance between 100% and 50% isoline (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Lateral Anlterior-Posterior Superior-Inferior</td>
<td>Lateral Anlterior-Posterior Superior-Inferior</td>
</tr>
<tr>
<td>PTV1</td>
<td>2.8 2.7 2.0</td>
<td>7.4 7.1 3.8</td>
</tr>
<tr>
<td>PTV2</td>
<td>4.0 3.9 1.6</td>
<td>10.5 14.5 3.6</td>
</tr>
<tr>
<td>PTV3</td>
<td>4.2 2.6 1.1</td>
<td>14.7 13.5 4.0</td>
</tr>
</tbody>
</table>

**Table 4. Dose fall-off for planning target volumes**

**Treatment delivery**

Following patient setup, XVI 3D cone beam CT was used for patient position verification prior to initiation of treatment. Translational and rotational corrections were made using the HexaPOD evo robotic treatment couch, which allows submillimeter patient alignment in 6° of freedom.

SRS was delivered by an Elekta Versa HD linear accelerator with the Agility MLC (collimator 0°, couch 0°) and a 6MV flattened beam, using a double 340° VMAT arc (gantry 190°-170°). The treatment consisted of 255 segments and 7309.85 MU. The total beam on time for the treatment of all 3 brain lesions using a double VMAT arc was just 12 minutes. This would have taken more than twice as long (25 minutes) if multiple isocenters had been used.

XVI 3D cone beam CT patient position verification was also performed post treatment to confirm the accuracy of the SRS treatment delivery.

The treatment was well tolerated by the patient and she was able to return home on the same day as treatment.
Outcome and follow up

Following SRS, the clinical symptoms of increased ICP decreased and the patient was comfortable to commence chemotherapy within one week.

Follow up MRI scans three months post treatment demonstrated lesion regression for all three brain metastases (Figure 5).

Figure 5.
Brain MRI scans
Discussion and conclusions

The standard protocol for brain metastases at Geetanjali Cancer Center

For solitary, limited (2–4) and multiple metastases, this institution follows ASTRO recommendations,¹ IRSA guidelines³ and additional published evidence⁴-⁶ for the use of SRS and WBRT (Figure 1). Recent ASTRO recommendations² state that adjuvant WBRT should not be added routinely to SRS for limited brain metastases due to the risk of diminished cognitive function, worse patient-reported fatigue and quality of life. Randomized studies have demonstrated no overall survival benefit with the addition of adjuvant WBRT to SRS in the management of selected patients with good performance status and solid tumor brain metastases.²

Radiosurgery alone, in cases where there are no additional metastases and greater than 6 months median life expectancy (less than 5% of the patient population at this institution), is only used in a trial setting at this center and does not form part of the standard protocol currently.

For multiple brain metastases with disseminated disease and less than 6 months life expectancy (greater than 95% of the patient population), patients are stratified according to the distance of the nearest lesion to the hippocampus. If the distance is greater than 4 mm, then hippocampus sparing WBRT is used, with or without SRS boost, in order to reduce the risk of cognitive decline. If the distance is less than 4 mm, then WBRT is used with or without SRS boost. A narrow margin (4 mm) around the hippocampus, used in hippocampus sparing WBRT, is achievable due to the increased modulation that can be delivered using Agility without compromising plan evaluation criteria. Furthermore, the positional accuracy of the HexaPOD patient positioning system (0.3 mm, 0.3°) provides confidence for accurate treatment delivery.
In the delivery of linac-based SRS to multiple targets in the brain there are a number of particular challenges.

**Accurate delineation**
Accurate delineation is essential for dose targeting precision and the sparing of normal tissue and critical structures in the brain. Any errors in SRS are critical due to the high doses delivered in a single fraction and the proximity of critical structures. For this reason, it is important to use the same slice thickness for the planning MRI and CT scans to minimize errors when the images are fused together during registration. This is essential for high precision procedures and allows accurate delineation of targets and critical structures.

**Patient immobilization & image guidance**
The second challenge for minimizing errors in linac-based SRS is robust patient immobilization. The options are to use frame-based or frameless immobilization for cranial radiotherapy. At the Geetanjali Cancer Center, a frameless immobilization protocol using the Fraxion cranial immobilization system with a thermoplastic mask is used. This method improves patient comfort, compared to frame-based immobilization, and provides sufficient rigid immobilization for cranial radiotherapy techniques when used in combination with accurate image guidance procedures.

For this patient, XVI CBCT imaging was performed to verify patient position before and after treatment delivery, and the HexaPOD robotic couch was used to align the patient with submillimeter accuracy prior to treatment initiation. Since the treatment time was less than 15 minutes, intrafraction imaging was not performed in this instance. Generally, if treatment time is greater than 15 minutes and/or if 2 VMAT arcs are used in treatment delivery, XVI intrafraction imaging is performed to ensure there is no patient movement during delivery.
Treatment delivery

Accuracy and speed are essential for SRS delivery. Dose targeting precision is ensured by accurate delineation and data input during the planning process, as already described. Accuracy in dose delivery is also enhanced by reducing beam-on time and, hence, further minimizing the risk of intrafraction motion. Previously, using multiple isocenters, treatment times were much longer. With the accurate dose placement and speed of delivery that can now be achieved using Monaco TPS and Versa HD (using VMAT and Agility), in addition to the possibility of using a single isocenter for multiple targets with Agility, three brain metastases were treated in 12 minutes. A single isocenter also has the benefit of reducing the number of MUs to the patient and reducing planning time compared to a multi-isocenter plan. The reduction in treatment time increases patient comfort and reduces the risk of errors caused by intrafraction motion.

The alternative treatment option for this patient would have been whole brain radiotherapy (WBRT). Linac-based SRS allowed this patient to benefit from faster treatment with reduced risk of neurocognitive decline and resulted in immediate subjective improvement of symptoms and lesion regression 3 months post treatment. The aim of this treatment was to minimize treatment time and to improve overall comfort for the patient, which was achieved. Where there is not access to Gamma Knife® radiosurgery, this method is an option for clinics, allowing SRS to be considered for more patients.

This single isocenter method has been shown to increase treatment planning and delivery efficiency, reduce treatment time and ensure efficient use of MU. The next step is to use flattening filter free (FFF) radiotherapy, which provides the opportunity to further increase efficiency by reducing treatment times for SRS deliveries. For this particular case, subsequent to treatment, the flattened beam plan was compared to an equivalent plan created using FFF 6MV beams. The FFF plan was delivered to a phantom using a double VMAT arc in just 4 minutes compared to 12 minutes for the flattened beam delivery. In addition, the FFF beam produced comparable dose coverage to the flattened beam plan. Planning efficiencies are also being investigated, particularly applying single arc VMAT plans where applicable. These approaches may be of added value for providing further treatment efficiencies and reduced beam on times in the treatment of multiple brain metastases using linac-based SRS.
References


Disclaimer

This case study is based on the experience and application of medical experts, and is intended as an illustration of an innovative use of Elekta solutions. It is not intended to promote or exclude any particular treatment approach to the management of a condition. Any such approach should be determined by a qualified medical practitioner.

It is important to note that radiation treatments, while usually beneficial, may also cause side effects that vary depending on the area being treated along with other medical circumstances. The most frequent side effects are typically temporary and may include, but are not limited to, skin redness and irritation, hair loss, respiratory, digestive, urinary or reproductive system irritation, rib, bone, joint or soft tissue (muscle) pain, fatigue, nausea and vomiting. In some patients, these side effects may be severe. Treatment sessions may also vary in frequency, complexity and duration. Finally, radiation treatments are not appropriate for all cancers, and their use along with the potential benefits and risks should be discussed before treatment.
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We help clinicians to improve patients’ lives through our forward-thinking treatment solutions and oncology informatics, creating focus where it matters to achieve better outcomes.