Role of Brachytherapy in Head and Neck Cancer:

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What is Brachytherapy?

The term ‘brachytherapy’ derived from greek word brachio (short)

A form of Radiotherapy that involves treatment with radioactive sources (usually sealed) in contact with or close to target tissue

2 distinct characteristics

- High dose of radiation to the adjacent target volume
- Limiting dose to the surrounding healthy tissues
History

"Actually Radiotherapy started in the form of Brachytherapy"

1898:
Marie & Pierre Curie isolated Radium and work on Brachytherapy started.
History

Robert Abbe (American surgeon):
First used Ra technique for treatment of cancer.

1920-30s: “Quimby System”

1934: “Paterson-Parker”
History of Brachytherapy

- **1950-60s**: Advent of mega voltage type tele-therapy machines had provided treatment option with non-invasive procedure; EBRT treatment.
- So, there was decline in progress of interstitial Brachytherapy.

*Brachytherapy was considered ‘lost art’.*
History

- **1964**: Paris system
  Bernard Pierquin et al. used Ir192 after-loading interstitial implant.

- **1980-90s**:
  HDR after-loading, computer planning & optimization came in use.

*New possibilities in Interstitial and intracavitary Brachytherapy with advantages of HDR after-loading & computer optimization.*
What is so special about Brachytherapy?

It is the best conformal radiotherapy.

Has been able to keep it’s identity in the face of new challenge.
Different applications of Brachytherapy in head and neck cancer

- **Implant**: Most common application sources placed directly into the target tissues as in oral cancer

- **Intraluminal**: single line source introduced into the lumen of a organ (ca nasopharynx)

- **Mould**: In cancer of lip ,BCC of face where sources placed over the skin or mucosal surface
Why Brachytherapy in Head & Neck Cancer?

1. Total Dose Higher to the tumour with normal tissue sparing (Better therapeutic ratio)

   - Better local control and less side effects
     (In head and neck cancers, failure is predominantly local or loco-regional.)

3. More accessible sites, as in tongue, cheek etc.

4. Overall treatment time smaller and treatment finished before the repopulation starts
Indications of Brachytherapy in H & N Cancer

- As a sole modality: Radical treatment in early Cancer (>3 cm and node- negative) tongue, cheek, floor of mouth

- Used with EBRT: To boost the dose at site of greatest risk of recurrence

- As salvage treatment: In recurrent tumours
Contraindications of Brachytherapy

- Diffuse and extensive growth
- Distant metastasis
- Node-positive disease
- Tumors very close to or involving bone
- Not suitable for anaesthesia
Sites in Head & Neck region suitable for implant

1. Ant 2/3 tongue
2. Lip
3. Buccal mucosa
4. Floor of mouth
5. Soft palate.
6. Base of tongue as boost.
7. Tonsil
Selection of patients for Primary Implant.

- Early disease, >3 cms and node-negative
- No distant metastasis.
- Site accessible, superficial and away from bone
- Pt willing
- Radical treatment intent.
- Suitable for anesthesia.
Steps of Implant.....

Interstitial Treatment

Implant of catheters
Role of Radiotherapist

dosimetry
Role of Physicist
Implant technique
Paterson-Parker system of Implant

- Pre loaded needles (cesium)
- Planar and volume implants
- Distribution of needles guided by the area and geometry of implant
- Crossing needles used often
- Needle position parallel and interneedle distance 1 cm usually
- >1.5 cm thick target needs double-plane implant
- Dose calculated at 0.5 cm from each needle
- Basically implant guided and dose prescription based on the Paterson-Parker table.
Implant technique

Paris System of Implant:
1. Radioactive sources parallel & arranged so that their centres lie in same plane; ‘central plane’.

2. Linear activity of each source wire uniform.

3. Radioactive sources are equidistant.

4. When multiple plane implant used, implants placed in such pattern that they maintain equilateral triangle or squares at central plane.
Implant technique

Implant - I

Implantation done following Paris System of implant

Excellent Implant

Implant - II

Needles not equidistant & not in same plane.

Sub-optimum Implant
Dosimetric Consideration
Dosimetric Consideration

- GTV
- Needles
- Ref Vol./PTV
- Basal dose.
- Central plane.
- High dose vol.
- Mean Central Dose
- Min Target Dose
**Definitions:**

**Reference volume** = 85% of basal dose iso-dose curve covering the volume.

**Central plane** = plane perpendicular to the direction of sources formed by the centers of each source.

**MCD** = mean basal dose at central plane.

**MTD** = minimum tumor dose at the periphery of CTV.

**Basal Dose** = dose at central plane mid point between two equidistant sources.

**Treated Volume** = volume covered by ref. iso-dose.

**High dose Vol** = \( V_{150} \) iso-dose volume.

**Low dose Vol** = \( V_{90} \) iso-dose volume.

**Dose Prescription** =
- Reference vol should cover the CTV adequately.
- No high dose or low dose volume within the CTV.
- less normal tissue irradiated.
Changes in Modern day Implant

- Computer isodose calculations
- Dwell-weight optimisation of single high-activity stepping source
- Utilisation of 3D image to define target volumes

Basically changed from implant guided to image guided dosimetry
Both implant: Rules of Paris system implant followed

But both the implants same?
Implant - A

PTV completely covered by Ref. Vol

Less normal tissue in Ref Vol.

High dose vol. minimal

**Implant:**
- Well Conformed.
- Excellent dose uniformity
Implant -B

- Not Conformed.
- Significantly high dose non-uniformity GTV coverage by Ref. Vol not adequate
- More normal tissue in Ref Vol.
- High dose vol. unacceptable.
- Photon energy higher.
- Source characteristic different.

**Implant:**
- Not Conformed.
- Significantly high dose non-uniformity
Optimal Implant.
Excellent conformity & Homogenicity.

Sub-optimal Implant.
Not well conformed & non-homogenous.

So, though implant rules are well maintained, both the implants are not same on dosimetric consideration.
So, quality of implant depend not only on how the needles are placed in the target volume, but also on the dosimetric characteristics of the source & relative position of catheters in relation to target volume.
Dosimetric definitions:

*For assessment of quality of implant*

- Homogenicity Index (HI)(ICRU 58): \(\frac{\text{MCD} \times 100}{\text{MTD}}\)
- High dose vol \((V_{150})\) & low dose vol \((V_{90})\)
- Coverage Index (CI): \(\frac{\text{TV}_{\text{ref}} \times 100}{\text{PTV}}\)
- Conformity Index (ICRU 62): \(\frac{\text{PTV} \times 100}{\text{TV}}\)
- DNR: \(\frac{V_{150}}{V_{\text{ref}}}\)
- CDVH.
- ODR.
- Conformity number (CN).
How planning for an implant done?
Physical examination & staging

Pre-implant assessment & planning for implant
Our Experience:

Interstitial Implant done.
• Catheters are placed according to Paris System rules.
• Number and position of catheters placed according to tumor volume.
• No bleeding point ensured.
• Catheter ends are fixed to prevent displacement.
TPS Planning & Dosimetric Consideration
Our Experience:

Tumor Volume defined
Our Experience:

Catheters reconstructed
Our Experience:

CTV defined
Our Experience:

Treatment Vol defined
• Iso-dose curves plotted.
• Homogenicity & Conformity factors taken care.
• Dose received by normal tissue taken care.
Dose considerations

Different dose rates used in brachytherapy:
(As per ICRU report 38)

- Low dose rate (LDR): 0.4-2.0 Gy/h
  (Cesium 137 needles, Iridium wires)
- Medium dose rate (MDR): 2-12 Gy/h
- High dose rate (HDR): > 12 Gy/h
  (Iridium 192, Cobalt 60)
Dose and number of fractions by HDR

- Varies according to site, T stage and treatment objective, whether alone or with EBRT
- The total HDR dose should be biologically equivalent to LDR
- To decide no. of fractions, L-Q model is used
- In mouth cancer, LDR dose of 60 Gy (in 6 days) is eq. to 45 Gy HDR, which may be given in 7-10 fractions
- Total duration of therapy (EBRT + HDR) should be kept within 8 wks, to minimize repopulation
- Interval between HDR fractions should be 6 hrs minimum, to allow repair of normal tissue
- Every institution has it’s own protocol considering the radiobiology, tumour details and other logistics as well
CUMULATIVE DVH OF TARGET

DVH_3: Cumulative DVH on target. State: Consistent.
CUMULATIVE DVH OF MANDIBLE
Our Experience:

Treatment given
FOLLOW UP: TONGUE
Our Experience:

POST IMPLANT
Our Experience:
Our Experience:

12 MONTH
Our Experience:

15 Month
Our Experience:

18 Month
**Local Control Rates in Oral Cancer at different stages by Surgery or RT alone**

<table>
<thead>
<tr>
<th>Oral tongue</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
</tr>
</thead>
<tbody>
<tr>
<td>RT Alone (Curie)</td>
<td>86%</td>
<td>80%</td>
<td>68%</td>
<td>NR</td>
</tr>
<tr>
<td>RT alone (Pernot et al)</td>
<td>93%</td>
<td>65%</td>
<td>49%</td>
<td>NR</td>
</tr>
<tr>
<td>RT alone (MD Anderson)</td>
<td>80-90</td>
<td>60-85</td>
<td>30-50</td>
<td>24-40</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Surgery Alone</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
</tr>
</thead>
<tbody>
<tr>
<td>(Memorial Hospital)</td>
<td>85%</td>
<td>77%</td>
<td>50%</td>
<td>NR</td>
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</table>

<table>
<thead>
<tr>
<th>Floor of Mouth (RT alone)</th>
<th>T1</th>
<th>T2</th>
<th>T3</th>
<th>T4</th>
</tr>
</thead>
<tbody>
<tr>
<td>(MD Anderson)</td>
<td>75-85</td>
<td>60-80</td>
<td>30-50</td>
<td>5-30</td>
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Brachytherapy as sole treatment for oral cavity cancers

<table>
<thead>
<tr>
<th>Author</th>
<th>Fx size</th>
<th>no.fr</th>
<th>Eq.dose</th>
<th>Pt.no</th>
<th>L.C</th>
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</thead>
<tbody>
<tr>
<td>Dixit et al</td>
<td>3 Gy</td>
<td>20</td>
<td>65 Gy</td>
<td>3</td>
<td>-</td>
</tr>
<tr>
<td>Lau et al</td>
<td>6.5 Gy</td>
<td>7</td>
<td>63 Gy</td>
<td>27</td>
<td>53%</td>
</tr>
<tr>
<td>Inoue et al</td>
<td>6 Gy</td>
<td>10</td>
<td>80 Gy</td>
<td>14</td>
<td>100%</td>
</tr>
<tr>
<td>Donath et al</td>
<td>4.5-5</td>
<td>10</td>
<td>54-63</td>
<td>13</td>
<td>90%</td>
</tr>
<tr>
<td>Leung et al</td>
<td>5.5-6</td>
<td>10</td>
<td>71-80</td>
<td>13</td>
<td>100%</td>
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</table>
# HDR Brachytherapy as boost to EBRT for oral cavity cancers

<table>
<thead>
<tr>
<th>Author</th>
<th>EBRT (Gy)</th>
<th>HDR dose (Gy)</th>
<th>No. Fr.</th>
<th>Eq. dose (Gy)</th>
<th>No. Pt</th>
<th>L.C</th>
<th>survival</th>
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</thead>
<tbody>
<tr>
<td>Yu et al</td>
<td>50</td>
<td>2.7</td>
<td>6</td>
<td>67</td>
<td>12</td>
<td>79%</td>
<td>45%</td>
</tr>
<tr>
<td>Dixit et al</td>
<td>40-48</td>
<td>3</td>
<td>7</td>
<td>63-71</td>
<td>18</td>
<td>80%</td>
<td>---</td>
</tr>
</tbody>
</table>
Modern day implant more image guided

Implants still follow the Paris system basically, but dosimetry more flexible

Geometric and dose-point optimisations main advantages of modern day implant, but no amount of optimisations can make up for a bad implant

Quality of an implant is now more defined and uniform

More clinical experience and long term results will help to optimize the system further
Thank You