



Accelerated Partial Breast Irradiation



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Partial breast irradiation: Attractive alternative

Accelerated RT: shorter treatment duration

APBI: Accelerated Partial Breast Irradiation



5-6 Weeks of RT

Whole breast



1 week of RT

Tumor bed with
adequate margin



BCT: Patient perspective



- Do not opt for BCT due to inability to stay away from home for 6-8 weeks
- Small percentage of women do not take RT after BCT (14-20%)
- Dependents on other family members
- Some of the patients are earning members to support their families
- Cannot stay away from home

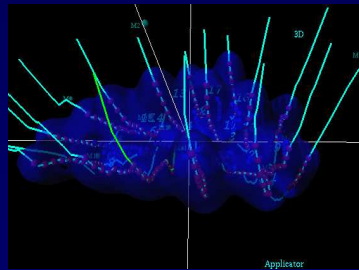
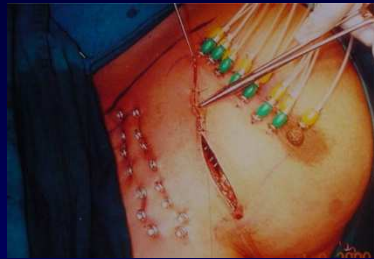




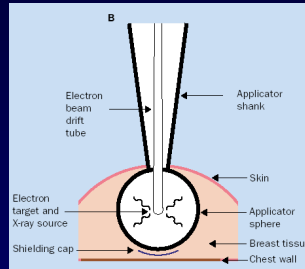
Clinico-pathological basis

- (70-90%) recurrences after whole breast RT in the tumour bed
- Pattern for site of recurrence same whether RT given or not (NSABP B06 trial)
- Very small percentage of the BCT patients recur outside tumour bed after whole breast RT
- Most of these outside recurrences are in fact New Breast Cancers
- Pathologically: multicentric foci seen away from the tumor bed
- But not all of them turn into cancers
- ? Need to treat whole breast in selected patient population

Methods of APBI



Interstitial brachytherapy



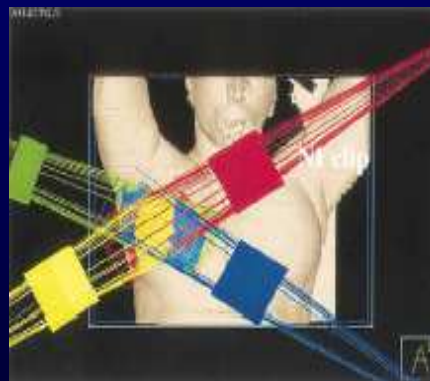
TARGIT



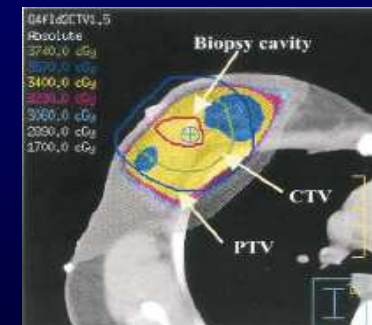
ELIOT



Mammosite



3DCRT



IMRT

Methods of APBI: Intraoperative X rays



TARGIT

Targeted intraoperative therapy

Source: 50KV Xray source

Technique: Intraoperative radiation after wide excision

Dose: 20Gy in 1 fraction at 1mm

Effective dose at 1cm: 5-7Gy

Advantage:

- simple technique

- sparing of normal tissues

Problems:

- Issues of penetration

- Adequacy of cavity wall dose ?

Encouraging early results

Methods of APBI: Intra-operative Electrons



Machine: Mobile linear accelerator

Electron energy: 3-10MeV

Technique: Wide excision

Placement of shield to protect chest wall

Reconstruction of the tumor bed

Dose: 21Gy at 90% isodose

Advantages: single fraction

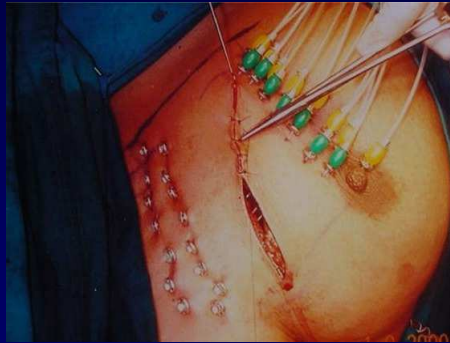
Problems: Issues of cavity wall coverage

Set up and expenses

Violation of surgical planes

Encouraging early results

Methods of APBI: Interstitial Brachytherapy



Brachytherapy



Oldest method

Large and encouraging data

Good target volume coverage with sparing of normal tissues

Brachytherapy Machines more common

Requires technical expertise

Methods of APBI: Mammosite



Mammosite

Balloon with single catheter

Dose: 34Gy/10 fraction BID

Advantage:

Ease of application

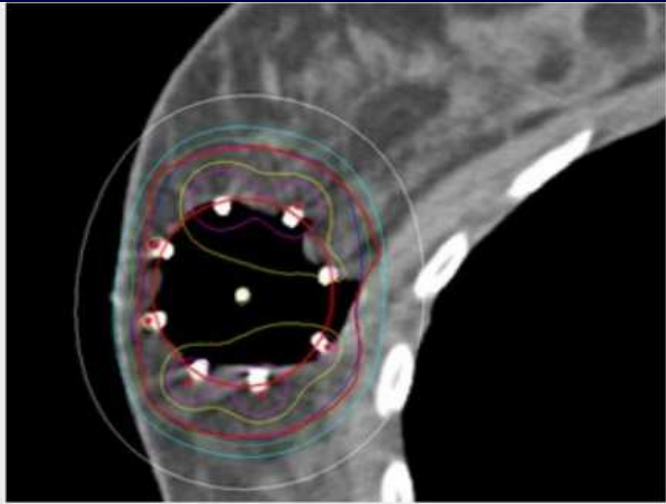
Problems:

High skin dose and telengectesia

Rib fractures

Problem in non-uniform cavities

Mutli-channel Catheters



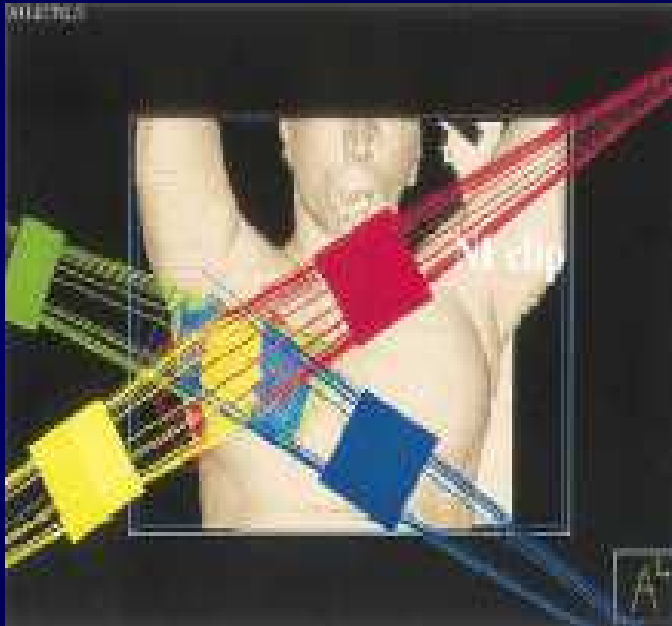
Mutlichannel Balloon based brachytherapy

Single balloon: to be inflated

Coverage better than Mammosite

Issues related to cavity coverage in irregularly shaped cavities

Methods of APBI: External Beam Radiation



Machine: Linear Accelerator

Technique: External Beam RT

3DCRT, IMRT, Tomotherapy

Advantages:

Good coverage of target

Good dose homogeneity

Problems:

Issues of movement with breathing

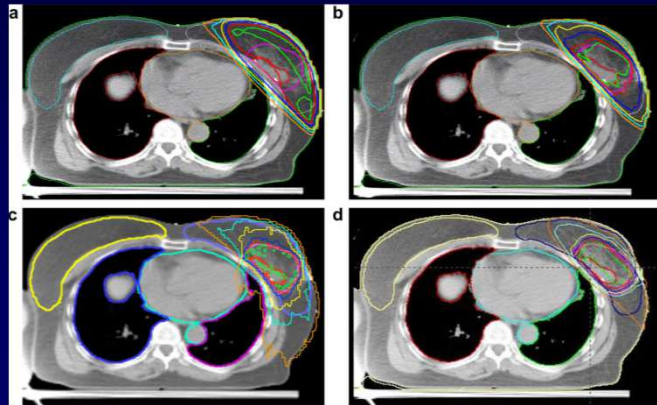
More margin

Higher integral dose-lungs, heart

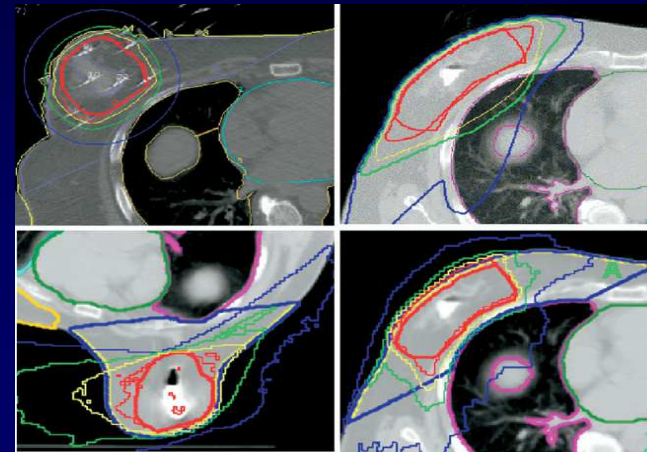
Comparison between the techniques

| | Interstitial Brachytherapy | 3DCRT/IMRT | Intraoperative electrons (ELIOT) | Intraoperative Xrays TARGIT | Mammosite |
|-----------------------------|--|-------------------------------------|--|--|--|
| Coverage of target volume | Variable | Best | Good | Good | Good |
| Thickness of target treated | 1-2cm | 2-2.5cm | 1-2.5cm | Dose prescribed at 1mm. At 10mm:5-7Gy | 1cm |
| Sparing of normal breast | good | least | good | best | good |
| Skin dose | Least | High | Least | Least (can shield) | Variable |
| Technical limitations | Axilla | Almost Nil | Axilla, brachial plexus, skin | Large cavities, irregular cavities | Large cavities, irregular cavities, close to skin, periphery |
| Drawbacks | Adequacy of target coverage Wider applicability | High dose to normal tissues, motion | Histopathology Wider applicability | Very limited depth of irradiation, cavity shape, size, no histopathology | Cavity shape and size Skin dose |

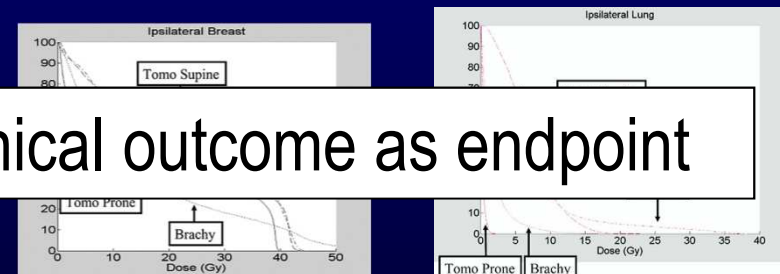
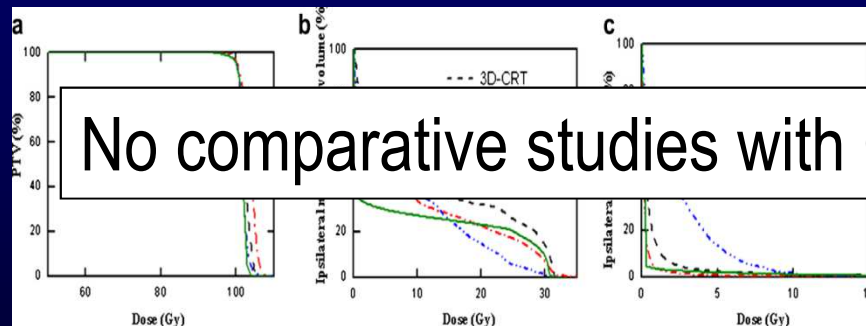
Comparison between the techniques



a. 3DCRT b: IMRT c: Helical Tomotherapy d: Proton therapy



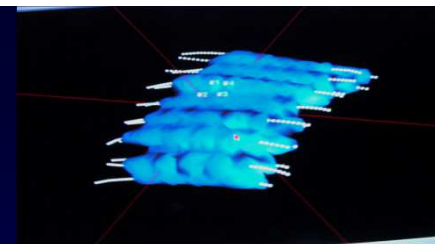
a. Brachytherapy b: 3DCRT c: Prone Tomo d: Supine Tomo



No comparative studies with Clinical outcome as endpoint



Selection Criteria for APBI



| Criteria | American Brachytherapy Society recommendation | American Society of Breast Surgeons recommendation |
|-------------|---|--|
| Age | 45 years or more | 50 years or more |
| Tumour size | $\leq 3\text{cm}$ | $\leq 2\text{cm}$ |
| Node | Negative | Negative |
| Histology | Infiltrating duct carcinoma (IDC) | IDC or DCIS |
| Margins | Microscopically negative | Microscopically negative ($>2\text{mm}$) |

Importance of patient selection

APBI studies in optimally selected patients

| Study | N | Median FU (yrs) | Local Rec % |
|--|-----|--------------------|-------------|
| Polgar (2009) NIO, Budapest | 45 | 12 | 8.9 |
| Johansson (2009) Orebro Medical Centre | 51 | 7.2 | 5.9 |
| King T (2000) Ochsner Clinic, New Orleans | 51 | 6.25 | 2 |
| Arthur DW (2008) RTOG 95-17 | 99 | 7 | 6.1 |
| Mark (2009) J Arrington Cancer Centre | 192 | 5.4 | 4.2 |
| Antonucci (2009) William Beaumont Hospital, Detroit | 199 | 9.6 | 5 |

APBI in suboptimally selected patients

| Institution APBI technique | No of patients (Median FU yrs) | Criticism | Breast Recurrence |
|--|--------------------------------------|---|----------------------|
| Christie Hospital RCT External Electrons 40Gy/8#/10days | 353 (8) | Lobular ca -15% Margin NK or+ve 19% Inadequate coverage | 25% |
| Guys Hospital LDR 55 Gy over 5 days | 27 (6) | Positive margins 55%, EIC+VE 40% | 37% |
| Uzsoki Hospital Budapest LDR 50Gy in 10-22 hrs | 70 (12) | Cut margin NK, single plane, unacceptable dose rate | 24% |
| London Regional Cancer Centre Ontario | 39 (7.5) | Av. Implant vol:30cc | 16% |
| Tufts New England | 33 (5) | 55% EIC | 6% |
| University of Kansas | 25 (4) | Inadequate LDR dose | 0% |

ASTRO Consensus statement :APBI outside clinical trial

| Factor | Suitable group | Cautionary | Unsuitable |
|---------------------|----------------------------|------------------|-------------------|
| Patient Factors | | | |
| Age | ≥60years | 50-59 | Age <50 years |
| Pathologic factors | | | |
| Tumor size | ≤2cm | 2.1-3 | >3cm |
| T stage | T1 | T0, T2 | T3,T4 |
| Margins | Negative (> 2 mm) | Close (<2mm) | Positive |
| Grade | Any | | |
| LVSI | No | Limited/focal | Present extensive |
| ER status | Positive | Negative | |
| Multicentricity | Unicentric only | | Present |
| Multifocality | Unifocal | | >3cm |
| Histology | Invasive ductal ,favorable | Invasive lobular | |
| Pure DCIS | Not allowed | ≤3cm | >3cm |
| EIC | Not allowed | ≤3cm | >3cm |
| Nodal factors | | | |
| N stage | pNo | | PN1, N2, N3 |
| Nodal surgery | SN Bx or ALND | | |
| Treatment factors | | | |
| Neoadjuvant therapy | Not allowed | | Used |

Similar guidelines given by GEC-ESTRO

Smith BD. IJROBP 2009;74:987-1001

TARGIT trial

Patients suitable for BCT

Age > 45 years, T size upto 3 cm,
unifocal tumors

BCT+ TARGIT

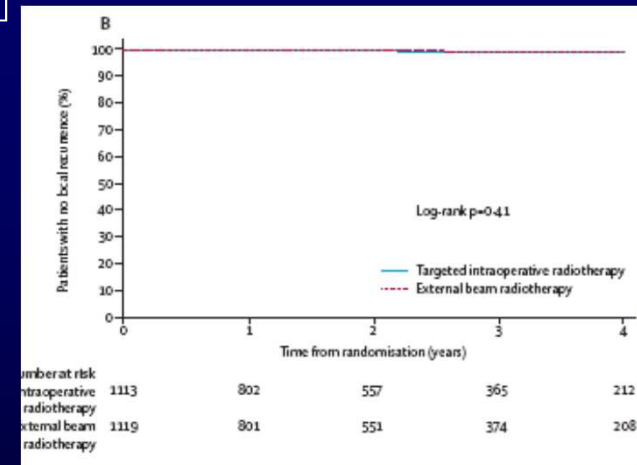
BCT+ External RT

N= 2232

Local recurrence rate at 4 years

TARGIT group: 1.2%

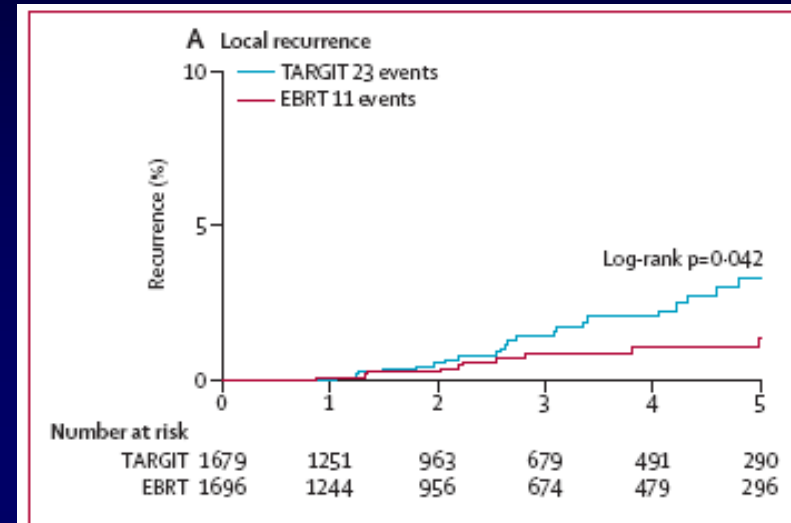
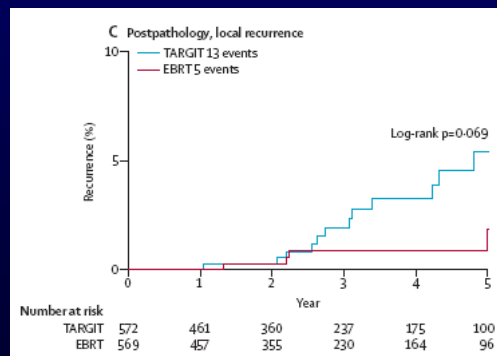
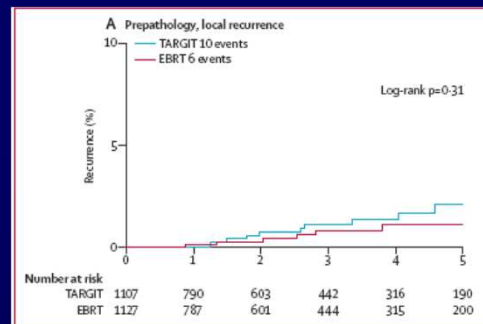
External RT group: 0.95%



TARGIT: 5 year Outcome

Local Recurrence in TARGIT arm: 3.3%

Local recurrence in EBRT arm: 1.1%



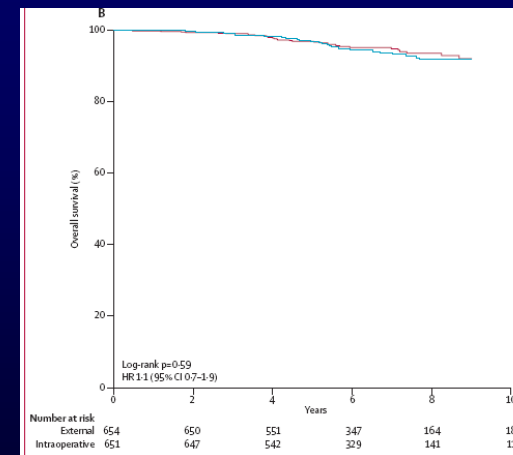
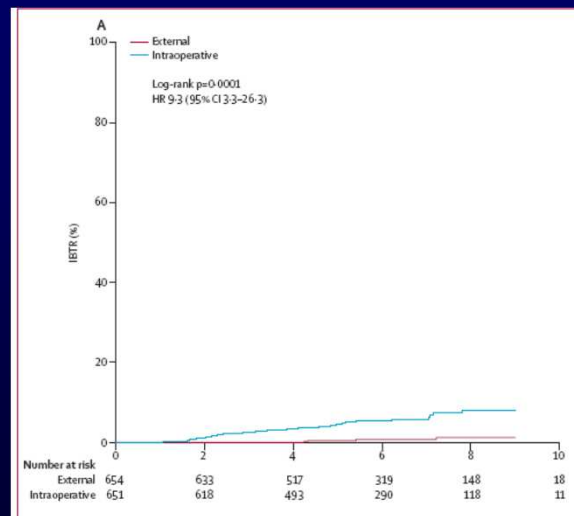
**TARGIT inferior to EBRT for
Local Control**

ELIOT (Intraoperative Electrons): Outcome

November 2000-December 2007
N=1306
T<2.5CM, Age >48 years

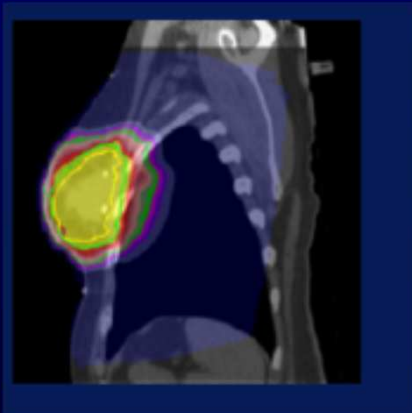
BCT+ Whole Breast RT
(60Gy) N=651

BCT+ ELIOT
(21Gy) N=655



3DCRT Technique: Outcome

- Prospective IRB approved study of Beamlet IMRT with deep inspiratory breath hold method.
- Dose: 38.5Gy in 10 fractions, 3.85Gy with bid regimen.
- 32 patients were enrolled
- With a median follow up of 2.5 years, 7 patients developed unacceptable cosmetic outcome.
- V50 and V100 volumes correlated with cosmetic outcome



Mammosite: 5 year outcome (phase II data)

- 1440 women
- Median FU: 53.7 months
- Median age: 65.5
- Median T size 1cm
- Node negative: 83.2%
- ER positive: 62%
- Grade I and II: 76.1%
- 5 year LR control rates: 96.2%
- Symptomatic seroma rates: 13%
- Excellent cosmetic outcome at 5 years: 90.6%
- ER negativity only strong factor affecting LR rates ($p=0.0022$)

Phase III data

Hungarian Randomized trial: 5 and 10 year results

1998-2004 (N=258)

T₁N_{0-1mi} breast cancer, low risk

Non lobular cancers, Clear margins, No EIC

Whole Breast RT(N=130)
50Gy/25#

Partial Breast Irradiation (N=128)

Interstitial brachytherapy (N=88)

Electrons (50Gy/25#) (N=40)

| | | |
|-----------|--------------|--------------|
| 5 yr LR | 3.4 % | 4.7 % |
| 5 yr OAS | 91.8 % | 94.6 % |
| Cosmesis | 62.9% | 77.6 % |
| 10 yr LR | 5.1% | 5.9% |
| 10 yr OAS | 82.1% | 79.7% |

Median FU: 66 months and 10.2 yrs respectively

Polgar C et al. Radiother Oncol 2013

Polgar C et al. IJROBP 2007; 69(3):694-702

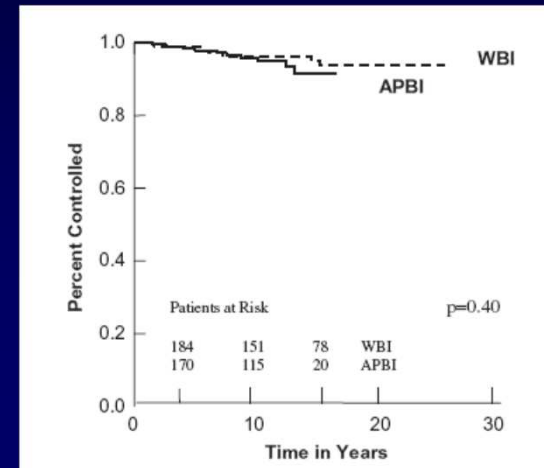
Phase II Data

12 year outcome of APBI: Match pair analysis

199 patients with interstitial brachytherapy

Matched with 199 women with whole breast RT

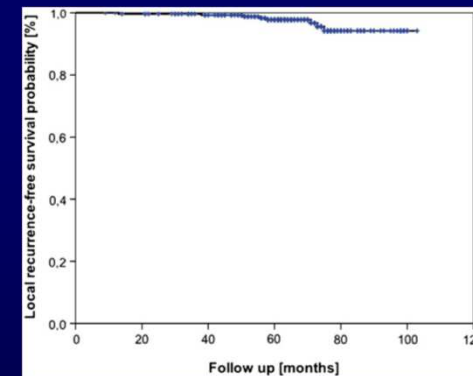
| | WBI (n = 199) | Interstitial APBI (n = 199) | p-value |
|--------------------------------|------------------|--------------------------------|---------|
| Age at diagnosis, mean (years) | 63.5 | 65.1 | 0.11 |
| Tumor size (mm) | 12.3 | 11.7 | 0.31 |
| ER+ | 85% | 86% | 0.85 |
| PR+ | 67.5% | 69.4% | 0.73 |
| Margins | | | 0.05* |
| Negative | 99.5% | 97.5% | |
| Positive | 0.5% | 0% | |
| Close | 0% | 2.5% | |
| T-Stage | | | 0.10 |
| T1 | 86.9% | 92.0% | |
| T2 | 12.6% | 8.0% | |
| T3 | 0.5% | 0% | |
| Lymph node status | | | <0.001* |
| Node negative | 88.4% | 88.4% | |
| Node positive | 2.0% | 11.6% | |
| Unknown | 9.5% | 0% | |
| Adjuvant hormonal therapy | 57.3% | 39.7% | <0.001* |
| Adjuvant chemotherapy | 3.5% | 12.6% | <0.001* |
| Follow up (years) | 14.0 | 10.4 | <0.001* |



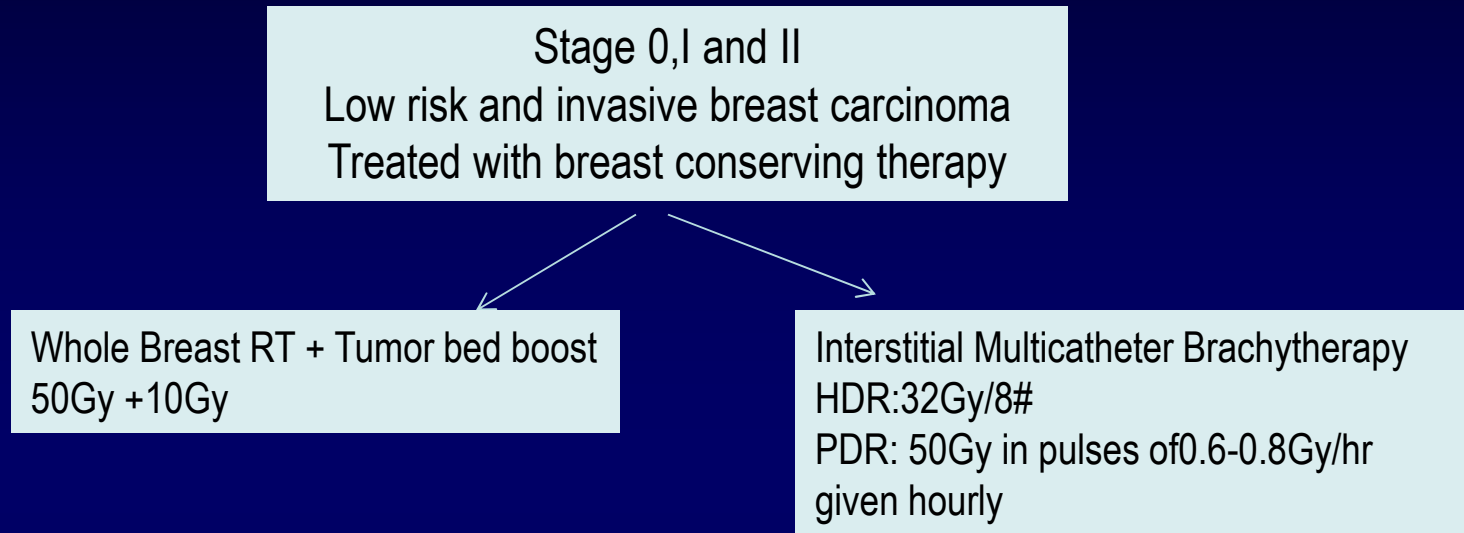
| 12 yr actuarial | WBI (%) | Interstitial APBI (%) | P value |
|-----------------|---------|-----------------------|---------|
| LR | 3.8 | 5 | 0.40 |
| RR | 0 | 1.1 | 0.15 |
| DFS | 87 | 91 | 0.30 |
| DM | 10.1 | 4.5 | 0.05 |
| OS | 78 | 71 | 0.06 |

German Austrian multicentric phase II trial

- Eligibility: Age > 35 years, T size <3cm, no lymph nodes, margins >2mm, hormone receptor +ve, histological grade I and II.
- N=274
- Median follow up 63 months
- Median Age: 60.5 years
- Median T size : 12 mm
- Chemotherapy: 6.9%
- 5 year local control rates: 98%
- 5 year DFS and OAS: 96.5% and 97% respectively



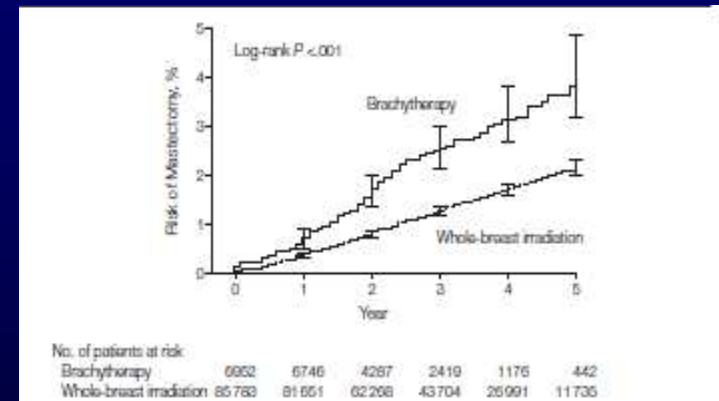
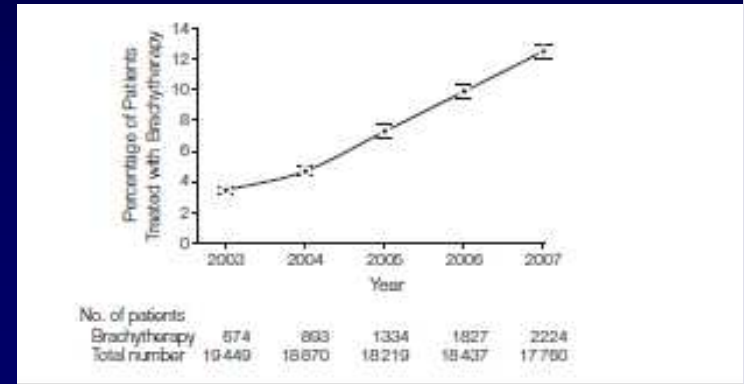
GEC-ESTRO Randomized trial of APBI



| | WBRT | APBI | P value |
|------------------------------|--------|--------|---------|
| 5 year Local Recurrence | 0.97% | 1.38% | 0.53 |
| 5 year disease free survival | 94.45% | 95.03% | 0.79 |
| 5 year overall survival | 95.5% | 97.25% | 0.11 |

Higher Complications with brachytherapy

- 92375 Women, medicare population
- Use of brachytherapy: 3.5% in 2003 to 12.5% in 2007
- 5 year Cumulative risk of mastectomy: 3.95% with brachytherapy and 2.18% with whole breast RT.
- Postoperative complications:
Brachytherapy: 26.5% WBRT: 16%
- Brachytherapy: Increased risk of breast pain, fat necrosis, rib necrosis

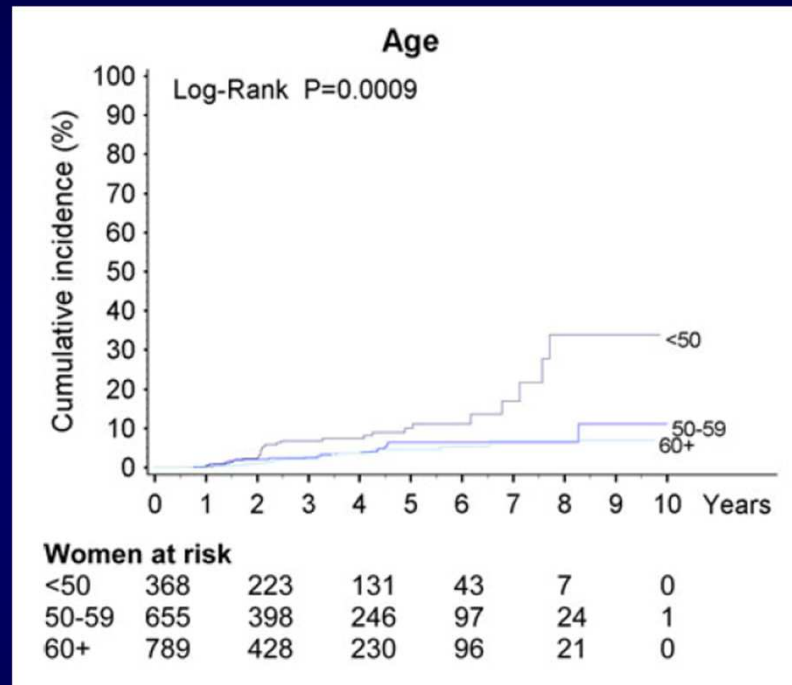


Brachytherapy: Mammosite in majority

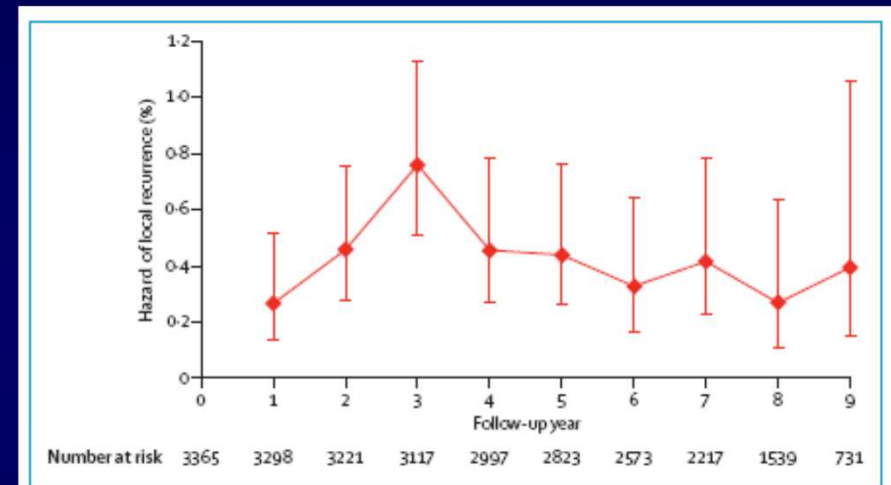
Other ongoing randomized trials

| Trial | Selection criteria | Technique in APBI arm | Target accrual and present status |
|---------------|--|--|--|
| NSABP, USA | Any age; <3cm DCIS or invasive ca with –ve margins and <4 nodes + | Interstitial or MammoSite HDR (34Gy/10#) or 3D CRT (38.5Gy/10#) | Target accrual-3000 patients >2000.Closed for low risk patients |
| Import Low | 50 years, pT > 2 cm, pN0 non-lobular, grade I or II, neg margins >2 mm | External RT: IMRT 40Gy/15# External RT: 36Gy/15# low risk, 40Gy/15# high risk | Target accrual 1935 Accrual completed |
| RAPID Ontario | >40 years, DCIS, <3cm, pN0 Nonlobular no BRCA1,2 | 3DCRT: 38.5Gy/10# | Target accrual 2128 Ongoing |

Importance of Long Term Follow up



Intraoperative electrons:
Outside trial



START trial data

Intraoperative Brachytherapy

W/E+ Axillary dissection



Confirmation of basic histopathological features on Frozen section



If suitable: Intraoperative placement of catheters in 2-4 planes



Radiotherapy planning X rays and CT scans on day 2/3



Treatment starts: day 3/4



Confirmation of final HPR before 5th fraction



Favorable: continue brachy



Unfavorable: convert to boost

Ext RT to be followed

APBI at TMH

Initiated in May 2000

Inclusion:

Age >40, T size upto 3cm, Node negative

No NACT, EIC -ve, C/M-ve

Till July 2016: 365 patients treated with APBI

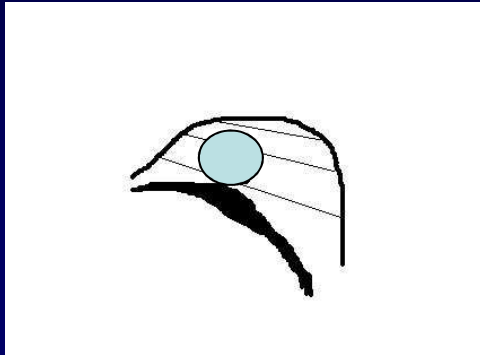
- Clinical examination, Mammography
- Brachytherapy done at the time of lumpectomy
- Pre-surgical assessment important
- Close collaboration with surgeon, pathologist, medical physicist



Lumpectomy cavity after wide excision
and axillary clearance

Placement of radio-opaque markers at four corners and centre of the cavity

Intra-operative Brachytherapy



Implant volume may appear larger than the treated volume



Marking of the planes on the skin



Insertion of needles in first plane



Insertion of second and third plane

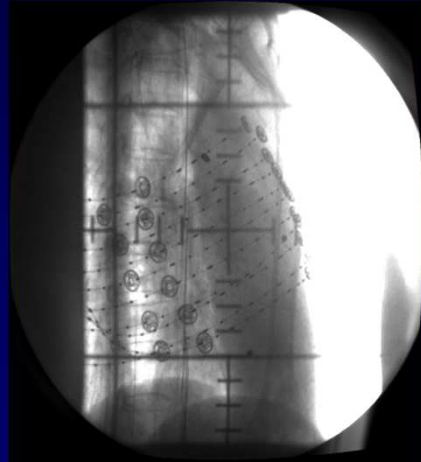
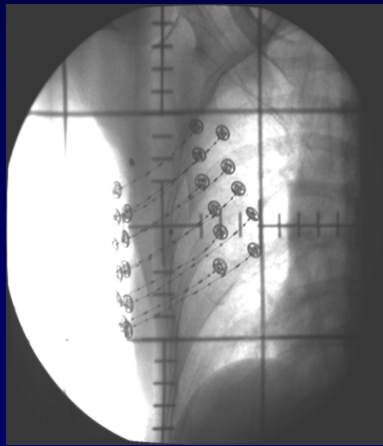


Replacement of needles with tubes

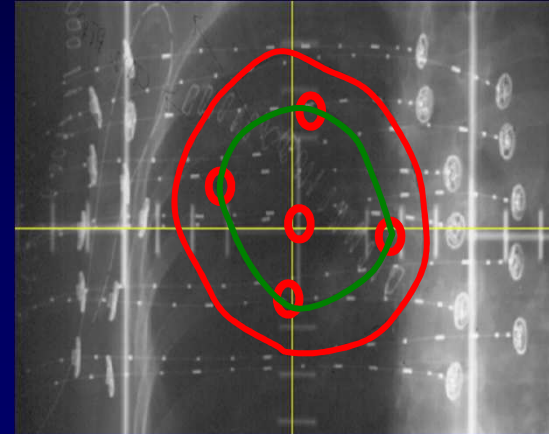
Post-operative Brachytherapy



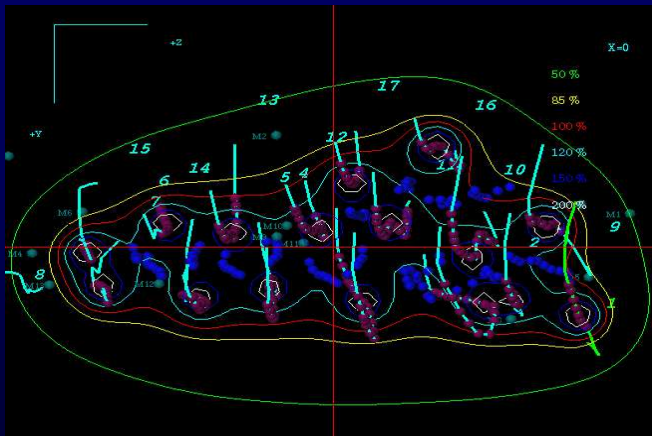
Brachytherapy Planning



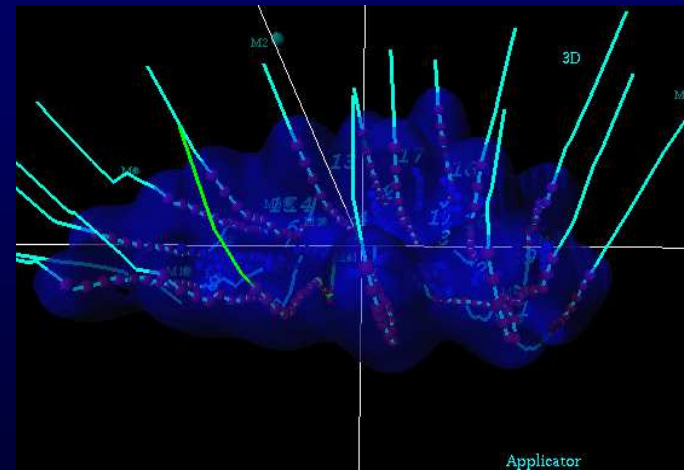
Orthogonal X rays



Identification of clips



Planning

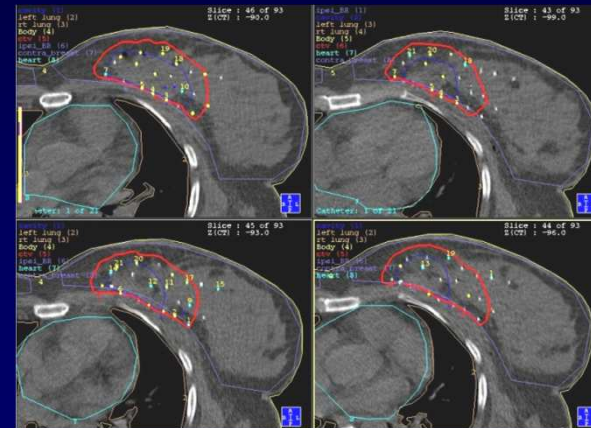


3D Dose distribution

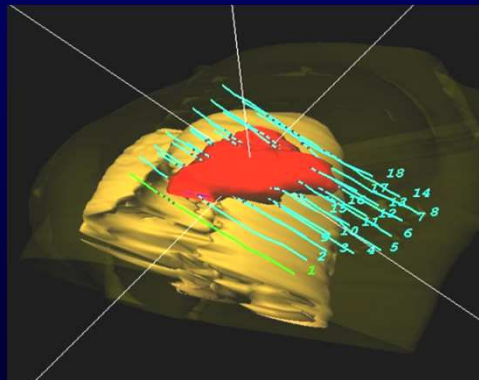
3D Brachytherapy planning



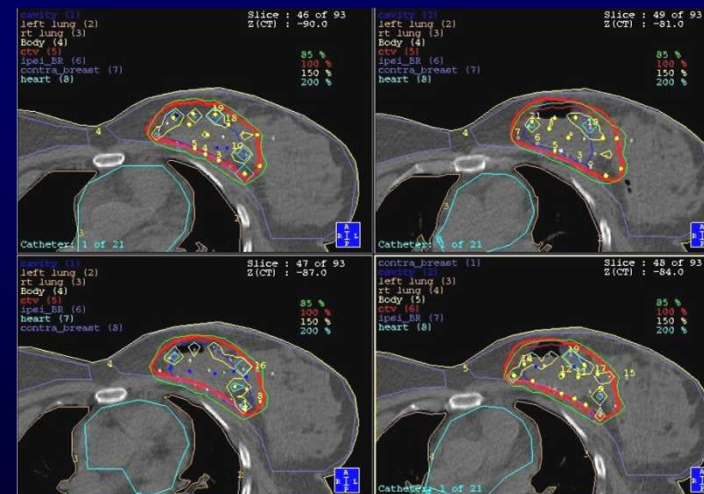
RT planning CT scan



Contouring



Determination of source loading



Slice by slice coverage evaluation

Treatment Delivery



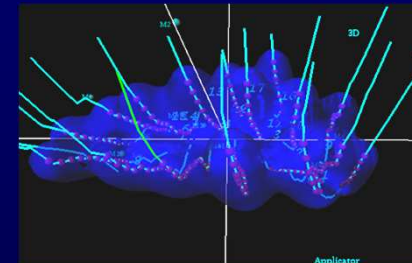
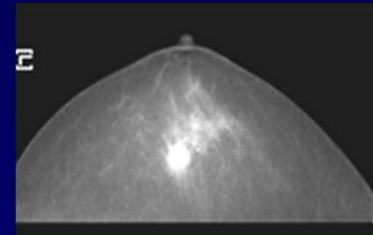
Dose: 34Gy in 10 fraction bid

Dose per fraction: 340cGy

APBI: TMH data: 2D Planning

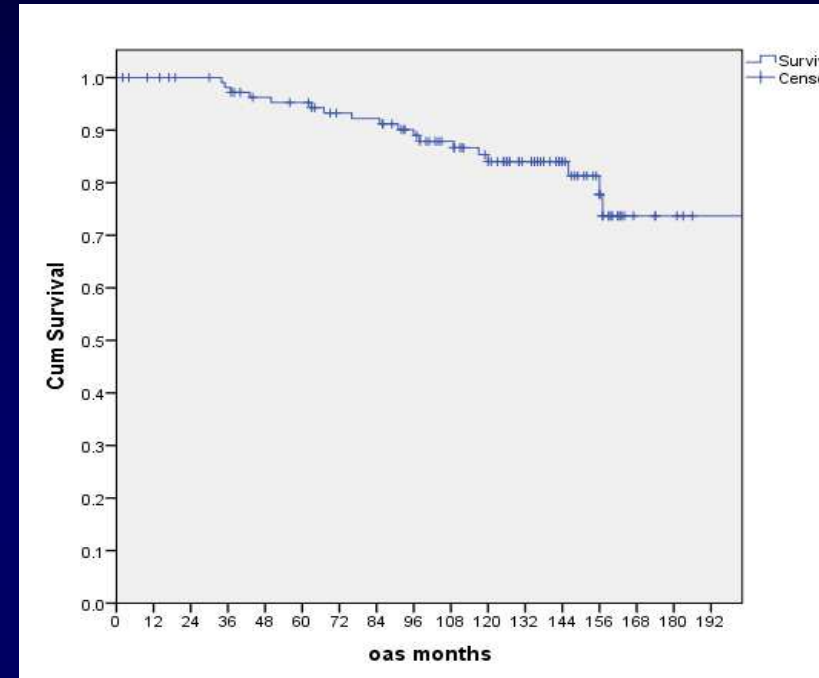
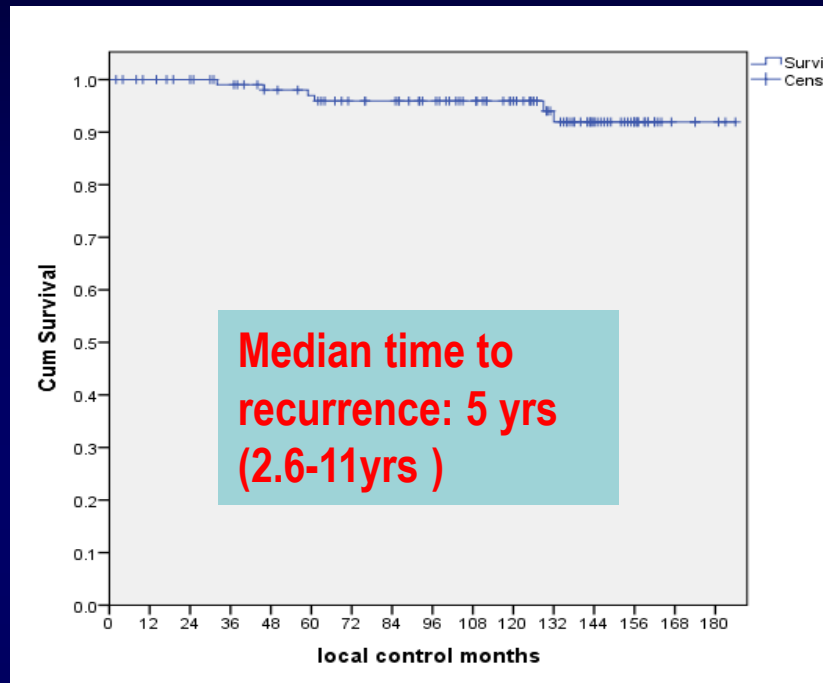
- May 2000- September 2005 (N=118) (X ray based)

- Median age: 56 years (30-78yrs)
- Median T size : 2cm
- IDC: 112 (97%)
- Grade III: 75 (65%)
- EIC positive: 8 (7%)
- Margin positive: 1 (1%)
- LVI: 13 (11%)
- Node positive: 12 (10%)
- ER positive: 62 (55%)
- Intra-op: 69 (60%)
- Chemotherapy: 55 (46%)



9 patients received WBRT
due to adverse prognostic factors

Clinical Outcome: > 10 year follow up



Median follow up 126 months

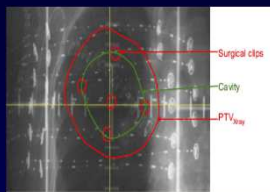
| | 5 yr | 10 yr |
|-----------------------|------|-------|
| Local Control | 97% | 96% |
| Disease free survival | 92% | 83% |
| Overall survival | 95% | 84% |

Analysis-July 16

Original Article

Dosimetric Comparison of Conventional Radiograph- and Three-dimensional Computed Tomography-based Planning using Dose Volume Indices for Partial Breast Intraoperative Implants

S. D. Sharma*, A. Budrukkar†, R. R. Upreti*, A. Munshi†, R. Jalali†, D. D. Deshpande*



18 patients-treated with APBI

| | P _{xray} | P _{CT} | P _{CT+graphical} | P value |
|-----------|-------------------|-----------------|---------------------------|--------------|
| CI Cavity | 0.80 | 0.82 | 0.92 | <0.001 (gr) |
| CI of PTV | 0.69 | 0.71 | 0.85 | <0.001 (gr) |
| DHI | 0.81 | 0.81 | 0.71 | <0.001 (gr) |
| OI | 0.041 | 0.047 | 0.087 | <0.0001 (gr) |
| EI | 44 | 25 | 30 | 0.013 (CT) |
| COIN | 0.48 | 0.58 | 0.68 | <0.001 (gr) |

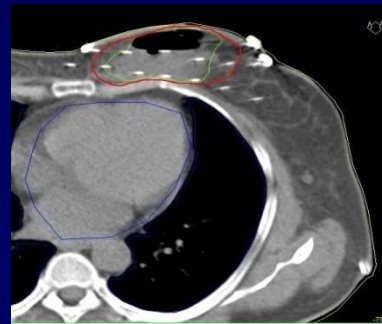
Conclusion: CT better than X ray for planning

APBI using 3D CT Based Brachytherapy

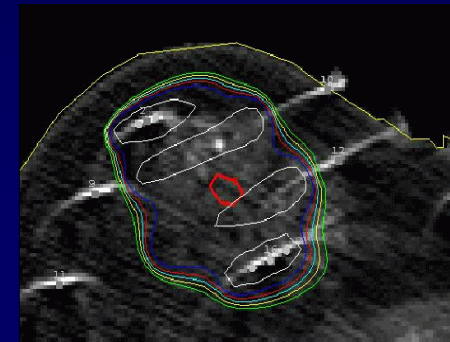
- Prospectively collected data: Between August 2005 to January 2013
- **Number: 140**



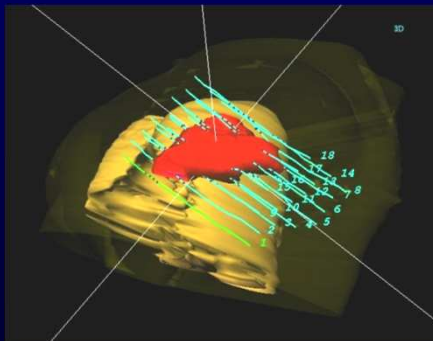
Planning CT scan



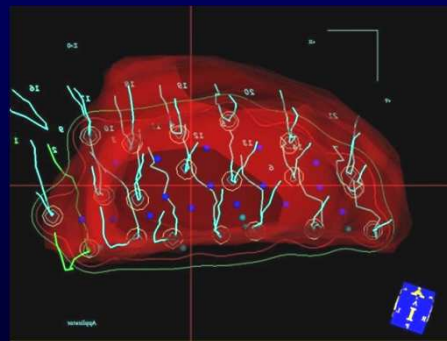
Contouring



Multiplanar reconstruction



Loading of sources



Dose points



3D Visualization



Partial breast brachytherapy

Clinical outcomes of prospectively treated 140 women with early stage breast cancer using accelerated partial breast irradiation with 3 dimensional computerized tomography based brachytherapy



Ashwini Budrukkar^{a,*}, Lavanya Gurram^a, Ritu Raj Upreti^b, Anusheel Munshi^a, Rakesh Jalali^a, Rajendra Badwe^c, Vani Parmar^c, Tanuja Shet^d, Sudeep Gupta^e, Tabassum Wadasadawala^a, Rajiv Sarin^a

^a Department of Radiation Oncology; ^b Department of Medical Physics; ^c Department of Surgical Oncology; ^d Department of Pathology; and ^e Department of Medical Oncology, Tata Memorial Hospital, Mumbai, India

- Median Age: 57 years (40-79)
- Postmenopausal: 109 (77.5%)
- Intra-operative brachytherapy: 80 (57%)
- Median T size: 2 cm (0.6-3.2cm)
- IDC: 140 (100%)
- Chemotherapy: 73 (52%)

Grade III: 115 (82%)

LVI: 11 (7.4%)

Margin positive: 1 (0.7%)

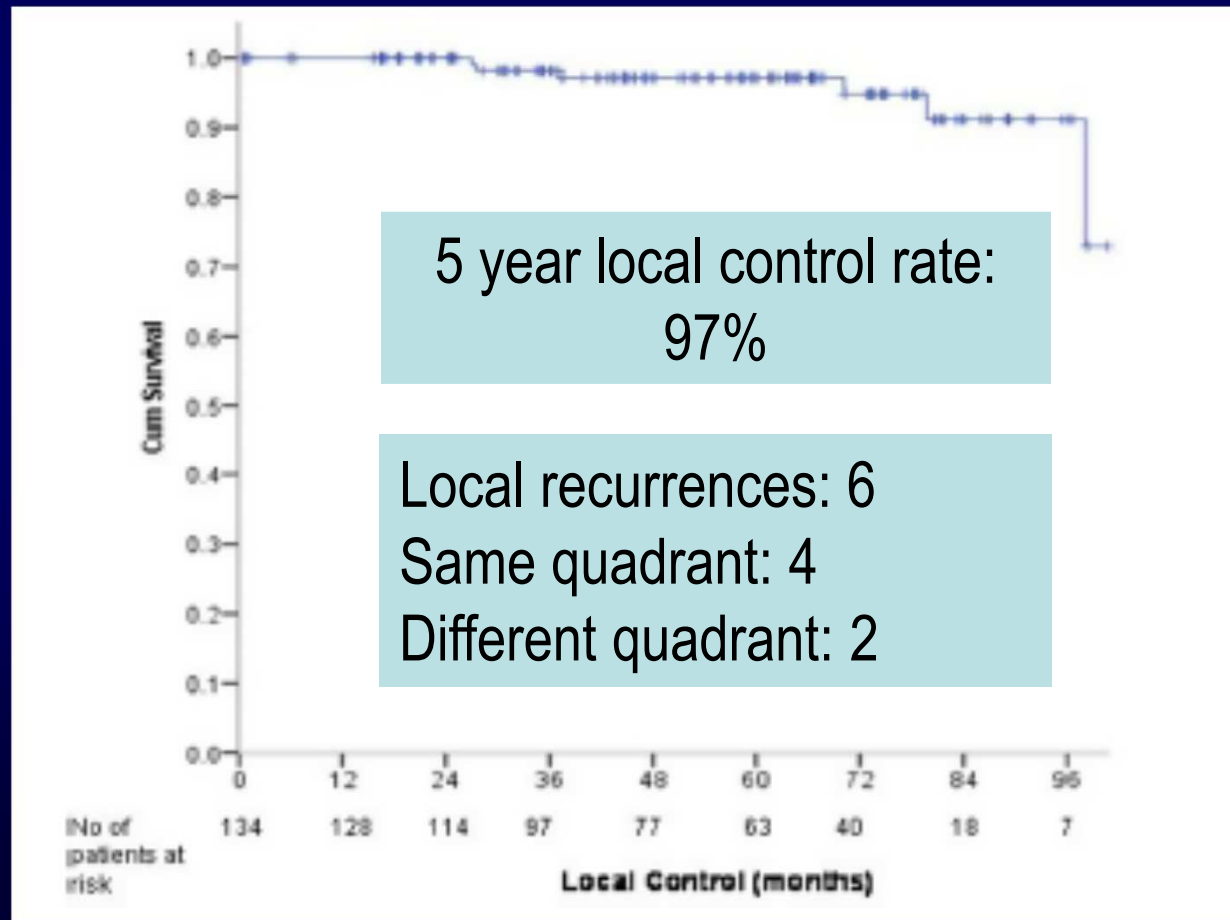
ER positive: 84 (60%)

Her2 positive: 23 (16%)



3DCT Based brachytherapy: Clinical Outcome

Median follow up : 60months (Range: 1-102months)



Median time to recurrence: 4.4 years
(2.2-6.5 years)

5 year OAS: 97%

Prognostic factors

| Factor | 5 yr Local control (%) | P value |
|---------------------------------|------------------------|---------|
| Age | | |
| <50 | 100 | 0.75 |
| ≥50 | 97 | |
| Pathological T size | | |
| ≤2 | 98.5 | 0.79 |
| >2 | 95 | |
| Grade | | |
| II | 100 | 0.34 |
| III | 98 | |
| Ductal carcinoma in situ | | |
| Yes | 96 | 0.25 |
| No | 98.2 | |
| Estrogen receptor status | | 0.16 |
| Positive | 100 | |
| Negative | 92.4 | |
| Her2 | | |
| Negative | 99 | 0.01 |
| Positive | 88 | |
| Vol 340 | | |
| ≤140 cc | 98 | 0.5 |
| >140cc | 100 | |
| Implant | | |
| Intra-op | 96 | 0.07 |
| Post-op | 100 | |

Cosmesis



Good to excellent cosmetic outcome: 77%



Late sequelae of APBI brachytherapy

Fat necrosis in women with early-stage breast cancer treated with accelerated partial breast irradiation (APBI) using interstitial brachytherapy

Ashwini Budrukkar^{a,*}, Vikas Jagtap^a, Seema Kembhavi^b, Anusheel Munshi^a, Rakesh Jalali^a, Tanuja Seth^c, Vani Parmar^d, Ritu Raj Upreti^e, Rajendra Badwe^d, Rajiv Sarin^a

^a Department of Radiation Oncology; ^b Department of Radiology; ^c Department of Pathology; ^d Department of Surgery; and ^e Department of Medical Physics, Tata Memorial Hospital, Mumbai, India

- 2000-2008; 170 women treated with APBI
- Median FU: 48 months
- 20 women developed fat necrosis
- Median time to development: 24 months
- **5 year actuarial fat necrosis rate: 18%**



Original Article

Quality of Life after Accelerated Partial Breast Irradiation in Early Breast Cancer: Matched Pair Analysis with Protracted Whole Breast Radiotherapy

T. Wadasadawala*, A. Budrukkar*, S. Chopra*, R. Badwe†, R. Hawaldar‡, V. Parmar‡, R. Jalali*, R. Sarin§

- EORTC QLQ & BR 23
- 48 patients-study period: May 2006-December 2006
- 23 APBI & 25 WBRT
- Median FU: 3 years
- **APBI better than WBRT**
 - QLQ C30
 - Social functioning (p=0.025)
 - Financial difficulties (p=0.019)
 - BR 23
 - Body Image (p=0.005)

APBI Team

- **Radiation Oncology**

- Rajiv Sarin
- Rakesh Jalali
- Ashwini Budrukkar
- Tabassum Wadasadawala
- Santam Chakraborty

- **Surgical Oncology**

- Rajendra Badwe
- Vani Parmar
- Nita Nair
- Shalaka Joshi

- **Medical Oncology**

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- Satish Kolhe
- Sudershan Kadam

- **Residents**

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- Lavanya Naidu
- Prakash Pandit
- Many more..