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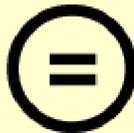
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**In vivo dosimetry and acute toxicity in
breast cancer patients undergoing
intraoperative radiotherapy as boost**

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In vivo dosimetry and acute toxicity in breast cancer patients undergoing intraoperative radiotherapy as boost

Directed by Professor Ik Jae Lee

The Master's Thesis

Submitted to the Department of Medicine,
The Graduate School of Yonsei University in partial
fulfillment of the requirements for the degree of
Master of Medical Science

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June 2017

This certifies that the Master's Thesis
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Jason Joon Bock Lee



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ABSTRACT

**In vivo dosimetry and acute toxicity in breast cancer patients
undergoing intraoperative radiotherapy as boost**

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(Directed by Professor Ik Jae Lee)

Purpose: To report the results of a correlation analysis of skin dose assessed by in vivo dosimetry and the incidence of acute toxicity. This is a phase 2 trial evaluating the feasibility of intraoperative radiotherapy (IORT) as a boost for breast cancer patients.

Materials and Methods: Eligible patients were treated with IORT of 20 Gy followed by whole breast irradiation (WBI) of 46 Gy. A total of 55 patients with a minimum follow-up of 1 month after WBI were evaluated. Optically stimulated luminescence dosimeter (OSLD) detected radiation dose delivered to the skin during IORT. Acute toxicity was recorded according to the Common Terminology Criteria for Adverse Events V4.0. Clinical parameters were correlated with seroma formation and

maximum skin dose.

Results: Median follow-up after IORT was 25.9 weeks (range, 12.7-50.3). Prior to WBI, only 1 patient developed acute toxicity. Following WBI, 30 patients experienced grade 1 skin toxicity and 3 patients had grade 2 skin toxicity. Skin dose during IORT exceeded 5 Gy in 2 patients: with grade 2 complications around the surgical scar in 1 patient who received 8.42 Gy. Breast volume on preoperative images ($p = 0.001$), ratio of applicator diameter and breast volume ($p = 0.002$), and distance between skin and tumor ($p = 0.003$) showed significant correlations with maximum skin dose.

Conclusion: IORT as a boost was well-tolerated among Korean women without severe acute complication. In vivo dosimetry with OSLD can help ensure safe delivery of IORT as a boost.

Key words: Boost with intraoperative radiotherapy, breast conserving surgery, acute toxicity, in vivo dosimetry

**In vivo dosimetry and acute toxicity in breast cancer patients
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I. INTRODUCTION

Whole breast irradiation (WBI) followed by tumor bed boost is standard radiation treatment in patients with breast cancer undergoing breast conserving surgery (BCS). Nevertheless, in selected patients with early breast cancer, partial breast irradiation (PBI) has shown acceptable local control rates with good cosmetic results.¹ Targeted intraoperative radiotherapy (TARGIT)-A trial used low energy x-rays (50 kV) generated from Intrabeam™ (Carl Zeiss, Meditec AG, Oberkochen, Germany) for delivering PBI during BCS and reported non-inferiority of intraoperative radiotherapy (IORT) compared to WBI using external beam radiation therapy (EBRT) in selected early breast cancer patients.² While American Society for Radiation Oncology (ASTRO) consensus guidelines recommend accelerated PBI for selected patients with early breast cancer,³ the feasibility of applying IORT as a boost to WBI in high-risk patients remains to be demonstrated. The use of

IORT as boost has been examined in retrospective studies^{4, 5} and a randomized comparison of these techniques with WBI followed by EBRT boost is currently underway in the TARGIT-B trial.

Intraoperative radiotherapy (IORT) delivers high dose to the tumor cavity and spares normal breast tissue; however, the radiation dose delivered to overlying skin and association with toxicity require close attention. Acute and late complications of IORT using IntrabeamTM include pain, seroma, wound infection, fibrosis, hyperpigmentation, edema of the breast, and ulceration.^{6, 7} One limitation of IORT is the lack of image-guided pretreatment plans, without which operators are unable to anticipate the radiation dose delivered to adjacent organs such as the overlying skin. Therefore, methods to assess radiation dose to the skin during IORT delivery (in vivo dosimetry) can be helpful in predicting the risk of developing skin toxicities and establishing proper inclusion criteria for future IORT applications. Since applying IORT as a boost has the potential to result in increased skin toxicity compared with IORT alone, additional safety evaluation is required for Korean patients whose breast volumes tend to be smaller than those of Western population. Since August 2014, a phase 2 trial investigating safety and feasibility of IORT used as a boost treatment in Korean patients requiring BCS has been underway at our institution. In the present report, we present the results of in vivo dosimetry and the correlation analysis of the factors associated with increased skin dose and acute toxicity.

II. MATERIALS AND METHODS

1. Study endpoints and eligibility

The primary endpoint was to provide an independent assessment of the delivered dose and consistency of the IORT technique through optically stimulated luminescent dosimeter (OSLD)-based in vivo dosimetry. The secondary endpoint was to determine clinical factors associated with increased risk of delivering a higher skin dose than intended. The phase 2 trial allowed enrollment of patients with histologically proven breast cancer undergoing BCS, aged 20 years or older, and with maximum tumor size less than 5 cm and Eastern Cooperative Oncology Group (ECOG) performance status of 1 or 2, but excluded patients who received neoadjuvant chemotherapy prior to BCS, or who required surgery for bilateral breast cancer. The study protocol conforms to the ethical guidelines of the 1975 Declaration of Helsinki, and was approved by our Institutional Review Board. All subjects provided informed consent prior to enrollment; all patients presented at a multidisciplinary tumor board at our institution.

2. IORT procedure

Two experienced surgeons were in charge of BCS. During BCS, a single fraction of 20 Gy was prescribed to the surface of the tumor cavity using the mobile 50-kV x-ray source (Intrabeam™) (Fig. 1). The x-ray source was attached to the robotic arm which maintained the stability of the source throughout the whole

process. Isotropy and the output of the unit were verified; the pre-IORT calibration process required by the system was performed prior to each treatment. Immediately after excision of the tumor, frozen sections were sent to the Department of Pathology for analysis to prevent positive resection margins after BCS. Re-excision was performed in cases of positive resection margins on frozen section analysis prior to IORT. A spherical applicator with an appropriate diameter (ranging from 1.5 to 5.0 cm in 0.5 cm increments) was selected according to the size of the tumor cavity, and the applicator was attached over the probe of the x-ray source. With a sterile sheath draped over the IORT device to prevent contamination, the spherical applicator was placed inside the tumor cavity, and a purse-string suture was used to pull the walls of the tumor cavity tightly against the applicator surface. The edges of the skin incision were everted so any part of skin was at least 1 cm away from the applicator surface to avoid excessive radiation exposure. When the skin-to-tumor distance was less than 1 cm, the portion of the skin overlying the tumor was excised to prevent suboptimal radiation dose delivery to the high risk area due to skin-edge eversion. Actual beam-on time after radiation site shielding was 20 to 30 minutes depending on applicator diameter. The surface of the tumor cavity received 20 Gy, while the radiation dose was attenuated to approximately 5 Gy at 1 cm depth. IORT was followed, with or without chemotherapy, by WBI of 46 Gy in 23 fractions.



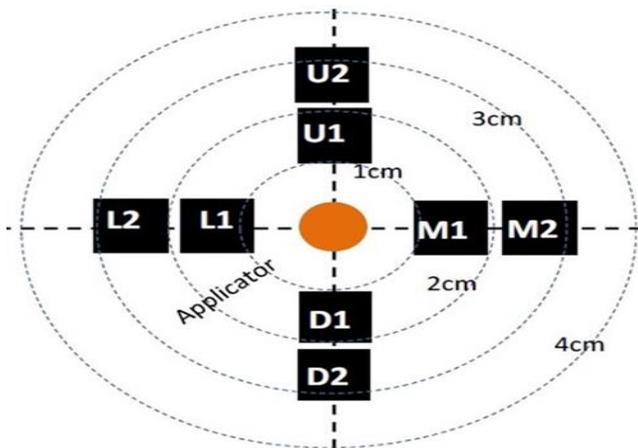
Figure 1. Intraoperative radiotherapy (IORT) device

3. In vivo dosimetry procedure

The OSLD used was InLight Nanodots (Landauer, Inc., Glenwood, IL, USA) which has a 5 mm diameter with thickness of 0.2 mm protected in a plastic casing. For each IORT measurement, 8 OSLD chips were annealed and attached to the skin, at 5 mm and 15 mm from the skin edge in the superior, inferior, medial and lateral locations (Fig. 2). After IORT, an InLight microStar® reader (Landauer, Inc., Glenwood, IL, USA) which utilizes a photomultiplier tube (PMT) to collect

the luminescence of the OSLD was used to determine the results of in vivo dosimetry. Each OSLD chip underwent measurements 3 times in the microStar® reader; the average PMT count was recorded.

A



B

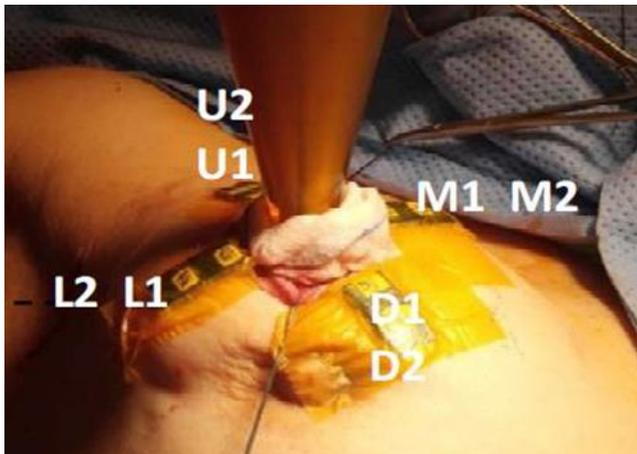


Figure 2. (A) Schematic arrangement of optically stimulated luminescent dosimeter (OSLD) and (B) example of OSLD arrangement in operating room

4. Toxicity evaluation breast volume measurement

Common Terminology Criteria for Adverse Events (CTCAE) Version 4.0 was used for toxicity assessment. Findings such as wound infection, edema, erythema and palpable seroma were documented 1 week and 1 month after IORT, at the end of WBI, and 1 month after WBI. Volumes of aspirated seroma were also documented. The ipsilateral breast of each patient on preoperative chest computed tomography images in supine position and breast magnetic resonance imaging (MRI) images in prone position was contoured according to the Radiation Therapy Oncology Group (RTOG) Breast Cancer Atlas for Radiation Therapy Planning. MIMVISTA 6.3.3. (MIM Software Inc. Cleveland, OH, USA) was used for contouring and calculating the preoperative breast volume of each patient on both imaging modalities.

5. Statistical analysis

Variables correlated with the maximum skin dose during IORT were determined using linear regression analysis; factors associated with seroma requiring aspiration 1 month after IORT were evaluated using logistic regression analysis. These variables included applicator diameter, skin-to-tumor distance in preoperative images, maximum tumor diameter, breast volume, volume of hemi-breast containing the tumor, ratio of applicator diameter and breast volume, and ratio of applicator volume and breast volume. The ratio of applicator diameter and breast volume was calculated as

$$\frac{\text{Applicator diameter (cm)}}{\text{Breast volume (cc)/1000}}$$

and the ratio of applicator volume and breast volume was calculated as

$$\frac{\text{Applicator volume (cc)}}{\text{Breast volume (cc)/1000}}$$

P values of <0.05 were considered significant; all statistical analyses were performed using IBM SPSS version 20.0 (SPSS Inc., Chicago, IL, USA).

III. RESULTS

1. Patients and treatment characteristics

From August 2014 until August 2015, IORT was considered in 124 patients; 82 patients actually underwent IORT and were enrolled in the phase 2 study. Among the 42 patients who did not undergo IORT, the reasons of IORT cancellation were either preoperative decisions in multidisciplinary conference, including presence of a huge primary mass or multicentric tumor nodules (n = 4); neoadjuvant chemotherapy (n = 3); definitive axillary, supraclavicular, or internal mammary lymph node metastasis (n = 3); and tumor close to the skin (n = 9); and intraoperative decisions including persistent positive resection margins (n = 9), conversion to total mastectomy (n = 8), and shape of tumor cavity unsuitable for IORT (n = 6). Among the 82 patients treated with IORT, 55 patients who underwent a minimum follow-up period of 1 month after WBI were included in the analysis. Follow-up after IORT ranged from 12.7 weeks to 50.3 weeks with a median follow-

up of 25.9 weeks. The median patient age was 52 years (range, 32-69), invasive ductal carcinoma was the most common type (69.1%), and tumors were most frequently located in the upper quadrant (87.3%). T stages according to the American Joint Committee on Cancer (AJCC) 7th edition were Tis in 7 patients, T1 in 40 patients and T2 in 8 patients; 4 patients were staged with N1 disease after operation with a single positive axillary lymph node. No patient had positive margins in frozen sections, but permanent pathology revealed positive margins in 3 patients. In these 3 patients, surgical re-excision was recommended; however, the patients refused. Sixteen patients received adjuvant chemotherapy and 37 patients received hormonal therapy. The median interval between IORT and of the initiation of EBRT was 5.7 weeks (range, 4.4-29.1). Patient, tumor, and treatment characteristics are summarized in Table 1. The median beam-on time was 18.15 minutes (range, 17.25-26.27); the diameters of applicators used were 2.5 cm (n = 3), 3.0 cm (n = 19), 3.5 cm (n = 28), and 4.0 cm (n = 5).

Table 1. Patient and treatment characteristics

Characteristic	No. of patients (%)
Age	
Median	52
Range	32-79
Pathology	
DCIS	7 (12.7)
IDC	38 (69.1)

Other	10 (18.1)
Site	
Left upper	25 (45.5)
Left lower	4 (7.2)
Right upper	23 (41.8)
Right lower	3 (5.5)
Tumor size (cm)	Median 1.4 (range 0.1-2.7)
≤1 cm	17 (30.9)
>1 cm	37 (67.3)
Unmeasurable	1 (1.8)
T stage	
Tis	7 (12.7)
T1	40 (72.7)
T2	8 (14.5)
N stage	
N0	51 (92.7)
N1	4 (7.3)
Number of positive lymph nodes	
0	51 (92.7)
1	4 (7.3)
Resection margin of permanent section	
Positive	3 (5.5)
Negative	52 (94.5)
Adjuvant chemotherapy	
Doxorubicin/Cyclophosphamide (AC)	15 (27.2)
AC + paclitaxel	1 (1.9)
No	39 (70.9)
Hormone therapy	

Yes	37 (67.3)
No	18 (32.7)
Interval between IORT & EBRT (months)	5.7 (4.4-29.1)

DCIS, ductal carcinoma in situ; IDC, invasive ductal carcinoma; IORT, intraoperative radiotherapy; EBRT, external beam radiation therapy

2. Toxicity

IORT followed by WBI was well tolerated. Although 1 patient experienced wound infection and required wound revision after BCS and IORT, no other patient experienced acute complications between IORT and the 1-month follow-up. Adverse skin reactions following WBI were observed in 33 (60%) patients, among whom 3 (5.5%) experienced grade 2 toxicity. More severe toxicity was not reported (Table 2). Fifty patients with seroma required aspiration at 1 week after IORT, 28 at 1 month after IORT, and 10 immediately after WBI.

Table 2. Toxicity (any grade) (n=55)

	1 week after IORT	1 month after IORT	Immediately after WBI	1 month after WBI
Skin Toxicity				
Erythema (grade 1) (n, %)	0 (0%)	0 (0%)	30 (54.5%)	2 (3.6%)
Erythema (grade 2) (n, %)	0 (0%)	0 (0%)	3 (5.4%)	0 (0%)
Wound infection (n, %)	1 (1.8%)	0 (0%)	0 (0%)	0 (0%)
Breast edema (n, %)	0 (0%)	0 (0%)	0 (0%)	0 (0%)

Seroma needing aspiration (n, %)	50 (90.9%)	28 (50.9%)	10 (18.2%)	n/a
Seroma (cc), mean (range)	15.4 (0-102)	5.5 (0-84)	1.6 (0-22)	n/a

3. *In vivo dosimetry*

At 5 mm from the skin edge, a maximum dose of 8.42 Gy was recorded, with a median dose of 1.76 Gy (range, 0.54–8.42) and a mean dose of 1.91 Gy. At 15 mm from the skin edge, a maximum dose of 6.58 Gy was recorded, with a median dose of 1.28 Gy (range, 0.44–6.58) and a mean dose of 1.41 Gy. Maximum dose recorded at 5 mm from skin edge was higher than maximum dose recorded at 15 mm from skin edge in every patient. As 20 Gy was prescribed to the applicator surface and the minimum applicator-to-skin distance was maintained at more than 1 cm, the expected maximum skin dose was approximately 5 Gy. In 2 patients, however, maximum doses of 5.33 Gy and 8.42 Gy were recorded; the breast volumes of these patients were 309 cc and 289 cc, respectively. The breast volume/applicator diameter ratios were 8.09 and 10.38. In both cases, the tumor was located close to the breast margin and the distance between skin and tumor, calculated from the dose distribution curve, was 1.3 cm for 5.33 Gy and 0.6 cm for 8.42. The patient with maximum dose of 8.42 Gy at 5 mm from the skin edge experienced grade 2 erythema at the spot where the maximum dose was measured (Fig. 3), although no skin reaction had been reported prior to WBI.

A



B



Figure 3. Patient with maximum dose of 8.42 Gy developed skin toxicity during EBRT. Photographs were taken (A) after 14 fractions of EBRT and (B) immediately after the completion of EBRT.

4. Risk factor analysis

The median breast volume was 567.8 cc (range, 157.2–1634.1) on MRI images in prone position and 511.4 cc (range, 230.1–1494.8) on CT images in supine position. The median volume of the tumor-bearing hemi-breast was 287.1 cc (range, 104.1–1015.4) on MRI and 287.2 cc (range, 112.3–969.2) on CT. In 6 (11%) patients, the tumor was located less than 1 cm from adjacent skin. Among 28

patients who presented with wound seroma needing aspiration 1 month after IORT, median and mean volume of seroma were 1.5 cc (range, 1.0-84.0) and 5.5 cc. According to our institution's criteria, 15 patients presented with considerable volume of seroma (> 5 cc). Analysis of factors associated with a considerable volume of seroma (> 5 cc) 1 month after IORT revealed that applicator diameter was a significant factor in both univariate ($p = 0.042$) and multivariate analyses ($p = 0.049$) (Table 3). Linear regression analysis showed a significant correlation between maximum skin dose and preoperative breast volume ($p < 0.001$), distance between skin and tumor on preoperative images ($p = 0.001$), and ratio of breast volume and applicator diameter ($p < 0.001$). Table 4 summarizes the results of linear regression analysis of factors associated with skin dose.

Table 3. Toxicity (Seroma > 5 cc)

	Univariate analysis		Multivariate analysis	
	95% CI	P value	95% CI	P value
Age	0.966-1.120	0.295	0.899-1.083	0.781
Applicator diameter	1.097-124.835	0.042	1.014-154.325	0.049
<div style="display: flex; align-items: center;"> <div style="margin-right: 10px;">≤3.5</div> <div style="margin-right: 10px;">>3.5</div> </div>				
Skin tumor distance	0.601-3.371	0.422	0.472-4.627	0.503
Maximum diameter on MRI (cm)	0.652-4.083	0.296	0.562-5.713	0.324

Diameter of applicator/ Volume of breast	0.657-1.151	0.329	0.615-1.299	0.557
Maximum skin dose	0.448-1.492	0.512	0.476-2.205	0.950

CI, confidence interval; MRI, magnetic resonance imaging

Table 4. Factors associated with skin dose

	R ²	P value
Skin-Tumor distance	0.184	0.001
Breast volume (CT)	0.254	<0.001
Breast volume (MRI)	0.274	<0.001
Diameter of applicator / Volume of breast (CT)	0.245	<0.001
Diameter of applicator / Volume of breast (MRI)	0.210	<0.001
Volume of applicator / Volume of breast (CT)	0.060	0.072
Volume of applicator / Volume of breast (MRI)	0.067	0.060

CT, computed tomography; MRI, magnetic resonance imaging

5. Receiver operating characteristic (ROC) analysis

Thirty-eight patients had a maximum skin dose larger than 2 Gy, while 17

patients had a maximum skin dose larger than 3 Gy. Median and mean skin dose were 2.26 Gy (range, 1.03-8.42) and 2.68 Gy. The inflection point of the maximum skin dose among the 55 patients was located between 2 Gy and 3 Gy. Therefore, patients recording maximum skin dose higher than 3 Gy can be regarded as larger than average group. ROC analysis was performed to distinguish useful parameters for predicting a maximum skin dose larger than 3 Gy. Among the risk factors mentioned above, the ratio of applicator diameter and breast volume was the best parameter for anticipating skin dose greater than 3 Gy. The area under the curve (AUC) was 0.820, and a threshold value of 7 resulted in the optimal sensitivity and specificity, 82.4% and 73.4%, respectively. The AUC of breast volume and the distance between skin and tumor were 0.165 and 0.267.

IV. DISCUSSION

IORT boost is known to offer several advantages over external beam boost in patients with breast cancer undergoing BCS. Immediate initiation of radiation after tumor removal may reduce the chances of tumor cell repopulation; direct placement of the applicator in the tumor cavity could reduce the possibility of targeting error that may occur due to daily setup errors and vague identification of the tumor bed on CT images acquired several weeks after surgery. Furthermore, dose distribution with steep dose fall-off allows increased dose to the tumor bed while reducing radiation to adjacent structures including skin and other organs such as the heart and lungs.^{8,9}

Although a few IORT-alone trials reported lower toxicity and survival rates comparable to those of WBI in low-risk patients with early breast cancer,^{2,7} higher local recurrence rates and lack of long-term follow-up of those studies prevent clinicians from moving to a single session of IORT from WBI.¹⁰⁻¹² On the other hand, IORT could be a method of delivering a boost dose in combination with WBI; its feasibility as a boost treatment has been examined retrospectively. Vaidya et al. reported that lumpectomy and TARGIT boost combined with WBI resulted in superior local tumor control compared to the results of the European Organization for Research and Treatment of Cancer (EORTC) boost trial or the Standardization of Breast Radiotherapy (START)-B trial.¹³ In addition, a single center experience at University Medical Centre Mannheim using low-kV X-ray IORT as a boost resulted in 5-year overall survival of 91.3% with 5-year local recurrence-free survival of 97.0%.⁶ An ongoing trial, TARGIT-B, is comparing IORT boost and EBRT boost in high-risk patients.

IntrabeamTM IORT requires inserting a spherical applicator in the tumor cavity and delivering a high radiation dose in a single fraction; thus, the toxicity risk is likely dependent on geometric variations such as applicator diameter, tumor location, and breast volume. However, no published studies clearly indicate exclusion criteria based on such geometric variables in the application of IntrabeamTM IORT.

As the breast volume of East Asian women is generally smaller than that of Western patients,¹⁴ it is imperative that the risk factors associated with increased toxicity are evaluated prior to the wide use of IntrabeamTM IORT in Korea. We

conducted an in vivo dosimetry study for patients undergoing Intrabeam™ IORT followed by WBI, and correlated geometrical variables of the breast with increased maximum skin dose.

Some researchers regard seroma as a natural consequence of PBI rather than a complication¹⁵ and have concluded that IORT is not associated with increased volume of seroma requiring treatment.¹⁶ Even though other factors than IORT boost, such as body mass index and age could contribute to the formation of seroma,⁵ careful observation is necessary after IORT, as wound seroma needing aspiration was reported more frequently in patients undergoing IORT than in patients undergoing EBRT in the TARGIT-A trial as well as in the report by Kraus-Tiefenbacher et al., in which 48% of patients had measurable seroma 1 month after IORT boost.^{2,5} Although neither of the study provided the optimum value for evaluation of seroma formation, we used 5 cc as a criteria considering mean volume of seroma 1 month after IORT boost was 5.5 cc. The present study showed that the applicator diameter was an independent predictor of significant seroma formation; however, whether or not the volume of seroma also depends on the high-dose irradiation during IORT remains to be determined.

Cosmetic outcome is an important factor that should be considered in patients with breast cancer. Cracco et al. reported comparable cosmetic outcomes between 81 patients treated with IORT and 105 patients treated with traditional EBRT.¹⁷ Further follow-up is required to assess the rate of late complications and cosmetic outcomes in Korean patients with breast cancer undergoing IORT boost followed by

WBI.

The novelty of this study is in determining selection criteria for high-risk patients using in vivo dosimetry during IORT. Recently, OSLD has been widely used in measuring multiple radiation dose types, such as heavy ion beams, photons, electrons, and protons. Sharma et al. successfully used OSLD in surface dose measurement for high dose rate (HDR) brachytherapy with minimal uncertainty.¹⁸ The data acquired from in vivo dosimetry were useful in determining factors related to higher skin dose, which may increase the risk of acute and late skin reactions. As we have illustrated, 2 patients in the present study received skin doses higher than 5 Gy despite efforts to keep adjacent skin at least 1 cm away from the applicator. Therefore, patients who have similar characteristics to these 2 patients, such as relatively small breast volume, tumor location close to the breast margin, and small skin-to-tumor distance, require careful attention. In the present study, preoperative breast volume, distance between skin and tumor on preoperative images, and ratio of breast volume and applicator diameter were found to be significant factors associated with higher skin dose; these factors must be considered before performing IORT. We are now using these pre-selection criteria in order to exclude patients with a high risk of skin toxicity at our institution.

The safety of IORT alone in the Asian population was reported by Sawaki et al., who evaluated the toxicity of IORT with 21 Gy in 5 Japanese patients with early breast cancer.¹⁹ The results of boost IORT in Asian patients are minimal; no institution has yet performed IORT for patients with breast cancer in Korea.

V. CONCLUSION

IORT boost was well tolerated among Korean patients with breast cancer, without severe acute toxicity. The data acquired in this study using in vivo dosimetry could be useful in selecting appropriate candidates for IORT boost in the future. The results of the phase 2 trial and the assessment of late toxicity will follow.

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ABSTRACT (IN KOREAN)

유방암 환자에서 수술 중 방사선치료를 boost로 사용하는 것에
대한 생체 내 선량 측정 및 급성 독성에 대한 연구

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목적: 유방암 환자에 있어 수술 중 방사선치료는 정상 유방 조직에 조사되는 방사선량을 줄이면서 종양이 있던 자리에 고선량을 조사할 수 있는 장점이 있다. 그러나 주변 피부 조직에 조사되는 선량과 그에 따른 부작용에 대해서는 주의 깊은 관찰이 필요하며 이번 2상 연구에서는 한국인 유방암 환자에 있어 수술 중 방사선치료를 boost로 사용하였을 때 주변 피부에 조사되는 선량과 그에 따른 급성 독성을 알아보하고자 한다.

대상 및 방법: 강남 세브란스병원에서 수술 중 방사선치료를 시행받은 55명의 환자를 대상으로 하였다. 대상 환자들은 수술 중에 20 Gy의 방사선치료를 시행 받고 이후 외부 방사선치료 46 Gy를 23회에 걸쳐 전체 유방에 시행받았다. 피부에 조사되는 선량을 측정하기 위해 광자극발광선량계를 사용하였고 CTCAE V4.0에 의거하여 급성 독성을 평가하였다.

결과: 추적 관찰 기간 중간값은 25.9주였고 수술 후 외부방사선치료 시행 전에 한 명의 환자에서 상처부위 감염으로 인한 급성 독성이 보고되었다. 또한 수술 중 방사선치료 시행 후

한 달째 추적관찰에서 흡인이 필요한 장액종은 28명의 환자에게서 보고되었다. 3.5cm 이상의 직경을 가진 장착기의 사용이 경우 5cc 이상의 장액종 형성과 관련 있는 것으로 나타났다 ($p=0.045$). 외부 방사선치료가 종료되는 시점에 총 30명의 환자가 1단계 독성을 경험했고 3명의 환자가 2단계 독성을 경험하였다. 광자극발광선량계를 사용하여 측정된 피부선량이 5 Gy를 넘었던 환자는 두 명이 있었고 8.42 Gy의 피부선량이 측정된 환자는 외부방사선치료 종료 시 흉터 주위에서 2단계 피부 독성을 보였다. 이번 연구에서는 수술 전 영상에서 측정된 유방 부피 ($p=0.001$), 장착기의 직경과 유방 부피의 비 ($p=0.002$), 피부와 종양 사이의 거리 ($p=0.003$)가 피부 선량과 관계 있는 요소로 밝혀졌다.

결론: 한국인 유방암 환자에 있어 수술 중 방사선치료는 심각한 특이 급성 독성 없이 진행되었다. 아울러 광자극발광선량계를 사용한 생체 내 선량 측정이 동반될 경우 안전하게 방사선을 조사할 수 있을 것으로 판단된다.

핵심 되는 말: 생체 내 선량 측정, 유방 보존술, 급성 독성, 수술 중 방사선치료