

Dosimetric Effects on skin of supine Immobilization Device in Intensity Modulated Radiation Therapy for breast cancer: a retrospective study

Ran Lv

Second Affiliated Hospital of Fujian Medical University

Guangyi Yang

Second Affiliated Hospital of Fujian Medical University

Yongzhi Huang

Second Affiliated Hospital of Fujian Medical University

Yanhong Wang (✉ 992415639@qq.com)

Second Affiliated Hospital of Fujian Medical University <https://orcid.org/0000-0001-8152-0574>

Research article

Keywords: Supine immobilization devices, Breast cancer, Intensity Modulated Radiation Therapy, Skin dose

Posted Date: September 30th, 2020

DOI: <https://doi.org/10.21203/rs.3.rs-66850/v1>

License:   This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

Version of Record: A version of this preprint was published at BMC Cancer on April 9th, 2021. See the published version at <https://doi.org/10.1186/s12885-021-08119-6>.

1 **Dosimetric Effects on skin of supine Immobilization Device**
2 **in Intensity Modulated Radiation Therapy for breast cancer:**
3 **a retrospective study**

4 **Ran Lv^{1*}, Guangyi Yang^{2*}, Yongzhi Huang³, Yanhong Wang[#]**

5 1, E-mail: 125375680@qq.com

6 2, E-mail: ygy.004@163.com

7 3, E-mail: 826477919@qq.com

8 * These authors contributed equally to this paper.

9 # Corresponding author: Yanhong Wang, Address: NO 950, Donghai Street, Fengze

10 District, Quanzhou, Fujian 362000, China. Telephone: +8618659731867 E-mail:

11 992415639@qq.com

12 **Abstract**

13 **Background:** Breast immobilization devices are commonly used in supine
14 breast radiotherapy while the dose perturbation effect is often overlooked in
15 Intensity Modulated Radiation Therapy for breast cancer (BC). This study is to assess
16 the dosimetric effect of supine immobilization devices on skin with a commercial
17 treatment planning system.

18 **Methods:**40 women with BC were divided into four groups according to the type of
19 primary surgery, group A and B corresponding to patients with left and right BC after
20 radical mastectomy, received a radiotherapy of 50Gy in 25 fractions, group C and D
21 corresponding to patients with left and right BC after breast-conservation surgery,

22 received a prescription of 40.05Gy in 15 fractions and tumor bed simultaneous
23 integrated boost to 45Gy. A 0.2 cm thick skin contour and two sets of body contours
24 were outlined for each patient. Dose calculations were conducted for the two sets of
25 contours using the same plan, the dose difference was assessed by comparing the
26 dose-volume histogram parameter results and by plan subtraction.

27 **Results:** The supine immobilization devices for BC caused a significantly increase in
28 the skin dose which may finally lead to skin toxicity. The mean dose increased by
29 approximately 0.5Gy and 0.45Gy in left (group A) and right (group B) BC after
30 radical mastectomy, 2.7Gy and 3.25Gy in left (group C) and right (group D) BC after
31 BCS; corresponding to group A, B, C, D, the V10 of skin increased 1.27%, 1.83%,
32 1.36%, 2.88% ; the V20 of skin increased 0.85%, 1.87%, 2.76%, 4.86%; the V30 of
33 skin increased 1.3%, 1.24%, 10.58%, 11.91%; the V40 of skin increased 1.29%,
34 0.65%, 10%, 10.51%. The dose encompassing of planning target volume, as well as
35 the HI and CI, showed little distinction between plan- and plan+.

36 **Conclusion:** The supine immobilization devices significantly increased the dose of
37 skin, especially for patients with BCS. The immobilization devices should be included
38 in the external contour to account for the dose attenuation and skin dose increment.

39 **Trial registration:** This is a retrospectively study and it has no intervention on human
40 health care, so this study was not registered.

41 **Key words:** Supine immobilization devices; Breast cancer; Intensity Modulated
42 Radiation Therapy; Skin dose

43

44 **Background**

45 Breast immobilization devices are commonly used in radiation oncology to
46 provide patient support and improve positional reproducibility during their
47 fractionated radiotherapy. In actual clinical practice, the beam attenuation and
48 build-up perturbations effect caused by the immobilization devices is often
49 overlooked because carbon fiber material are widely used in immobilization
50 devices and are believed to be radio translucent for mega-voltage photons(1).
51 However, the density of carbon fiber is not equivalent to air, attenuation and scattering
52 can occur when the radiation beam pass cross these immobilization systems(2, 3).
53 Previous researches have indicated that immobilization devices used in radiotherapy
54 reduced tumor dose, increased skin dose (bolus effect) and altered dose
55 distribution(4-6). A. De Puyseleyr and his colleagues have reported that irradiating
56 through the carbon fiber immobilization device for prone breast radiotherapy resulted
57 in a considerable beam attenuation (range from 5.33% to 7.57%) and degradation of
58 skin sparing(4). For Chinese BC patients, due to small and compact glands, supine
59 positioning is still the most common approach and has multiple advantages, such as
60 methodological simplicity, comfort and accurate, reproducible positioning, and
61 reduction mean dose to heart(7, 8).

62 When compared with conventional wedge-based breast radiotherapy,
63 intensity-modulated radiotherapy is capable of delivering highly conformal and
64 homogenous dose distribution to targets and, herein, significantly decrease clinical
65 toxicities such as dermatitis and edema(9-11), which, in turn, substantially increased

66 the adoption of Intensity Modulated Radiation Therapy (IMRT) during breast
67 radiotherapy(12, 13). However, the increased beams and monitor unit (MU) have an
68 increased propensity to deliver radiation beam through the immobilization devices,
69 and, then, radiation immobilization device attenuation happened, ultimately
70 compromising the target coverage and organs-at-risk (OARs) protection(14). But no
71 study has yet assessed the dosimetric effects of supine breast immobilization devices
72 on the delivered doses to the target volume and OARs for breast IMRT. In this study,
73 we quantified the dosimetric effect of supine immobilization device by comparing the
74 dose distribution calculated with the breast immobilization to that calculated without
75 it, and investigated the potential skin sparing for BC patients achievable with 6 MV
76 photon beams in IMRT plans.

77

78 **Methods**

79 **Patient data and setup**

80 Forty women with BC were randomly selected and enrolled in this study; they
81 were divided into four groups according to the type of primary surgery and the lesion
82 location. Patients with left or right BC receiving radical mastectomy were assigned
83 into group A or B respectively. In the same way, patients with left or right BC
84 receiving breast-conservation surgery (BCS) were divided into group C or D
85 respectively. The patients' age ranged from 32 to 65 years, with a median age of 47
86 years.

87 **Simulation**

88 All patients were simulated in the head first, supine position using the carbon
89 fiber breast bracket (Klarity Inc, Guangzhou, China) for body immobilization. The
90 supporting board inclined at a certain angle ($7^{\circ}/12^{\circ}/17^{\circ}/23^{\circ}$ for choose) to assure that
91 the sternum was horizontal. The head was positioned straight on a circle sponge head
92 support, with the chin slightly upwards, avoiding skin folds at the lower neck. Both
93 arms rose over the head using a pair of arm-support to expose the breast adequately,
94 and a knee support to prevent the body sliding down. A thermoplastic film (electron
95 density 0.3~0.7, thickness 2.4mm) (Klarity Medical Products, Newark OH) was
96 custom molded over the chest and attached to the bracket by a plastic batten (electron
97 density 1~1.1). Computed tomography (CT) image with a 3-mm-slice was performed
98 using a large aperture CT Simulation scanner (Brilliance, Philips Medical System,
99 Amsterdam, Netherlands) (Fig.1). The scan range was from the first cervical vertebra
100 to 2 cm below breast ruffle. The obtained simulation CT images were transferred to
101 the treatment planning system (TPS, Monaco V5.11, Elekta AB, Stockholm, Sweden)
102 for target and OAR delineation and formulate treatment planning.

103 **Regions of interest**

104 Regions of interest were delineated on CT images with the CT data set as soft
105 tissue (window 600, level 40) by experienced oncologists according to
106 recommendation of international Commission on Radiation Units and Measurements
107 Reports 83 (ICRU)(15) and China Society for Radiation Oncology (CSTRO)
108 consensus(16). For patients underwent breast-conservation surgery, the clinical target
109 volumes (CTVs) include CTV breast, which includes all mammary glandular tissue,

110 and CTV boost, which is defined as the tumor bed including clips and seroma plus 5
111 mm's margin in all directions without exceeding the CTV breast. The corresponding
112 planning target volumes (PTVs) were generated by uniformly expanding 5mm from
113 CTVs respectively. In view of smaller mammary gland of eastern female, the ventral
114 border is designed 3mm below the skin surface(16). For patients with modified
115 radical mastectomy, CTV include chest wall and regional lymph nodes, the ventral
116 border is next to the skin surface(16). Organs at risk (OARs) including heart, left
117 anterior descending artery (LAD), left ventricle (LV), contralateral breast, ipsilateral
118 lung, and contralateral lung were countered for left-side BC, and liver was countered
119 especially for right-side BC. In this study, to assess the surface dose variance from
120 immobilization devices in TPS, the skin contour was especially delineated in the
121 treatment region with 2mm thickness below the skin surface for each patient
122 [Fig.1](17).

123 As a commercial treatment planning system, an external structure, which should
124 contain the materials involving in calculation, must be defined for calculating the dose
125 distributions. In this study, two sets external body contours were created for each
126 patient: one set included only the patient body without immobilization devices, and
127 the other set included the patient external body contours and the whole breast
128 immobilization devices.

129 **Treatment planning and dose calculation**

130 The prescription doses to PTV boost and PTV breast were 45Gy and 40.05Gy,
131 respectively, with a total of 15 fractions in case of BCS. And, in the mastectomy

132 situation, a prescription of 50Gy in 25 fractions was given. All patients were planned
133 on Monaco TPS with the energy of 6MV, using the dynamic inverse IMRT technique.
134 Multi-beam IMRT employs three groups of similarly-opposed lateral field spaced
135 through a 290° -150° sector for left-side tumors and 200° -60° sector for right-side
136 tumors around the target volume, which includes the breast/chest wall and regional
137 nodes, as indicated in Fig.1. 0° field was added when periclavicular node region was
138 included. A 0.5cm bolus was added to the surface skin in the treatment region for
139 patients with radical mastectomy for compensating the build-up effect of X-ray; on
140 the contrary, bolus was avoided for patients with breast-conserving surgery for
141 improving the cosmetic outcome. The optimization was accomplished by Monaco's
142 build-in Monte Carlo (MC) algorithm combined with Dynamic Multi-Leaf Collimator
143 (DMLC) technology.

144 For each patient, two IMRT plans were generated. The IMRT plans without
145 immobilization devices taking into calculation were recorded as Plan-. Under the
146 same irradiation constraints, the dose distribution was recalculated with the external
147 body contour containing the immobilization device and this plan was recorded as
148 Plan+.

149 **Statistical Analysis**

150 Dose-volume histogram (DVH) was the popular method to evaluate the dose
151 coverage of PTVs and OARs. For the PTVs, the parameters were the mean dose
152 (Dmean), the homogeneity index (HI) and the conformity index (CI). HI and CI were
153 respectively calculated as follow(18, 19).

154
$$HI = \frac{D_{5\%}}{D_{95\%}} \quad (1)$$

155
$$CI = \frac{TV1}{TV} * \frac{TV1}{VR1} \quad (2)$$

156 In formula (1), $D_{5\%}$ and $D_{95\%}$ were the doses received by 5% and 95% of the
 157 volume of Region of Interest (ROI), respectively. The HI value closer to 1 indicates
 158 the better uniformity of dose distribution in target volume. In formula (2), TV1 is the
 159 volume of target that receives the prescription dose and TV is Target Volume. VR1 is
 160 the Total Volume within prescription isodose curve. The CI value is between 0 and 1,
 161 and higher CI value indicates better dose conformity. For the OARs, the average dose
 162 D_{mean} , sometimes the maximum dose D_{max} and the dose-volume were calculated.

163 For each patient, the dosimetric effect caused by the immobilization devices was
 164 calculated by plan subtraction in the TPS. \bar{D} represents the average of parameters
 165 difference between Plan+ and Plan- as in formula 3, and $\bar{D}\%$ represents the average of
 166 relative difference between Plan+ and Plan-.

167
$$\bar{D} = \sum_1^{10} [(plan+) - (plan-)] / 10 \quad (3)$$

168
$$\bar{D}\% = \sum_1^{10} \{ [(plan_n+) - (plan_n-)] * 100 / (plan_n+) \} / 10 \quad (4)$$

169 The Statistical Package for Social Sciences V22.0 software (SPSS Inc., Chicago,
 170 IL, USA) was used to analyze all data. The Wilcoxon matched-paired signed-rank test
 171 was used to evaluate the significance of the observed differences between the plan+
 172 and plan-. The differences were considered statistically significant when $p < 0.05$.

173 **Results**

174 The comparison of dosimetric difference between Plan- and Plan+ for BC
 175 patients receiving radical mastectomy are presented in Table 1 and 2. The parameters

176 (Coverage Index, Dmean, D_{2%}, and CI) of PTV have little difference, except for HI of
177 left-side cancer ($\bar{D} = -0.006$) and D_{98%} of right-side cancer ($\bar{D} = 0.38$ Gy) with
178 indistinctive difference. Due to bolus effect of breast immobilization devices, the
179 mean dose and relative volume receiving 10, 20, 30, and 40Gy of skin were
180 significantly increased for Plan+ (\bar{D} and $\bar{D}\%$ were 0.45Gy and 1.11%, 1.83% and
181 2.01%, 1.87% and 2.27%, 1.24% and 1.56% and 0.65 and 0.96% for right breast
182 cancer, and 0.50 Gy and 1.25%, 1.27% and 1.37%, 0.85% and 1.00%, 1.30% and
183 1.67% and 1.29% and 1.99% for left breast cancer, respectively, all $p < 0.05$, Fig. 3A).
184 However, there was no statistically significant difference in other OARs.

185 For patients with breast-conserving surgery, dosimetric effects of immobilization
186 devices were calculated between Plan- and Plan+ too. As showed in Table 3 and 4
187 and Fig. 2B, plans calculation with breast immobilization devices had higher mean
188 skin dose ($\bar{D}\% = 10.21 \pm 2.95\%$ for right breast cancer and $9.07 \pm 1.60\%$ for left lesion)
189 and more volume of skin receiving interested dose (10-40Gy) radiation ($\bar{D}\%$ were
190 3.17%, 5.84%, 16.49% and 51.63% for right breast cancer and 1.65%, 3.57%,
191 15.85% and 51.86% for left breast lesion, Fig. 3B), which again displayed the bolus
192 effect of immobilization devices. For left-side BC patients, the mean dose and relative
193 irradiation volume of interested dose of left lung were decreased with little clinically
194 significance in Plan+ (\bar{D} were -0.21Gy for Dmean, -1.18% for V₅, -0.98% for V₁₀,
195 -0.47% for V₂₀, -0.28% for V₃₀, respectively). When it came to planning target
196 volume, the coverage index, HI and CI were also altered with statistical but not
197 clinical significance. As for the other OARs which far away from planning target

198 volume, such as contra-lung, heart, LV, LAD, liver, showed indistinctive difference
199 between two plans.

200 **Dose difference distribution map (Plan+ - Plan-)**

201 The dose difference distribution was calculated by Plan+ subtracted Plan-. As
202 shown in Fig. 2A, the blue to red gradient represented different absolute dose values
203 ranging from -5Gy to 5Gy. The build-up effect and radiation scattering caused by the
204 immobilization devices dramatically altered the dose distribution. The skin dose was
205 observably increased in irradiation region when the immobilization devices were
206 taking into calculation. In other word, the skin dose was underestimated by
207 approximately 6Gy if the immobilization devices were not included in the external
208 contour. Dose in other regions including Lung-L and PTV was decreased, which is
209 similar to DVH and data comparison results.

210 **Discussion**

211 Patient immobilization devices have become an important tool to guarantee
212 accurate delivery of highly conformal dose distribution(20). As the materials used in
213 immobilization devices are not completely X-ray transmission and can cause
214 attenuation of delivering dose, the dosimetric effects of immobilization devices
215 should be included in dose calculations(14, 21) . In addition, the bolus-effect cannot
216 be avoided either. When the beams, especially the posterior oblique orientations ones,
217 passing through the immobilization devices involved in the treatments, the
218 unexpected skin does can be increased and the dosimitric effect would be modified
219 finally(5, 22).

220 Beam attenuation from the couch, additional inserts and immobilization devices
221 can cause a misrepresentation of the actual dose delivered to the PTV, a deviation by
222 more than the recommended 3% to 5% accuracy range were reported by Olch(20). Li
223 Chen found the attenuation of the head and neck immobilization devices, which
224 reduce dose coverage rate (reduced by from 1.51% to 9.92%) and average dose
225 (reduced by from 0.93% to 1.92%) of planning target volumes in nasopharyngeal
226 carcinoma(21). Alyssa Olson assessed dose variance from immobilization devices in
227 VMAT head and neck treatment planning, and found that plan calculated without
228 immobilization devices was problematic with compromising V95, D100 and PTV
229 coverage(14). However, in our study, there was no clinically important effect of
230 supine breast immobilization devices on dosimetric parameters of PTV and PGTV,
231 with less than 3% deviation. The potential reason may lie in that all radiation beam
232 does not pass through the couch in our study. Puyssseleyn *et al.* measured the
233 dosimetric impact of a prone breast immobilization device and found that beam
234 attenuation amounted to 7.6% (6 MV X-ray) for beam pass through the couch
235 top-base plate combination and almost 5% beam attenuation for beam traversing the
236 couch-top(4). Then, there are less than 3% beam attenuation happened when beam
237 passing through base plate only, which is similar to our study.

238 Radiation-induced skin toxicity (RIST) is one of the predominant side effects for
239 BC patients, and deserves consideration as severe skin toxicity can lead to cession of
240 treatment and cosmetic changes. Francesco *et al.* stated that breast skin receiving dose
241 $\cong 30\text{Gy}$ is the most predictive parameters of acute RIST(23). While Tsair-Fwu

242 showed that skin receiving a dose $>35\text{Gy}$ (V_{35}) was the most significant dosimetric
243 predictor associated with radiation dermatitis grade 2+ toxicity(24). In our study, a
244 bolus effect of immobilization devices has been observed as the skin mean dose and
245 volume of receiving 10-40Gy are significantly increased in Plan+. This effect is more
246 obvious for patients after BCS. And V_{30} seems to be the most sensitive parameter
247 except for patients with right breast cancer and receiving radical mastectomy.
248 Currently, there is no standard of practice to include immobilization devices within
249 body contour, then actual skin dose was underestimated, which in turn induced more
250 and severer dermatitis. Take all above, we suggest the immobilization devices should
251 be included in dose calculations, and skin of breast region should be delineated as an
252 organ at risk and a dose-volume constraint for skin should be defined whenever
253 possible.

254 Although the positive results of this study, it has its limitations yet, a larger
255 patient population, different TPS or calculation algorithms, dosimetry techniques and
256 dose measurements are needed for further study.

257 **Conclusions**

258 In this study, dosimetric effects on skin of supine immobilization devices for BC
259 were calculated and evaluated in IMRT. The data shows the skin dose was
260 significantly increased, especially for patients with BCS, both the $V_{30\%}$ and $V_{40\%}$ of
261 skin increased sharply more than 10%. This research will remind radiation
262 practitioners to pay attention to the skin dose caused by the immobilization devices
263 and seek solution to remove the negative effects finally, continued research is

264 imperative. We prefer to include the immobilization devices within external body
265 contour and account for the skin dose increment in the TPS calculation.

266

267 **Declarations**

268 **Ethics approval and consent to participate**

269 This study was approved by the Second Affiliated Hospital of Fujian Medical
270 University Ethic Committee [2020(371)]. As the data are anonymous and no
271 intervention was happened in the treatment of patient, the Second Affiliated Hospital
272 of Fujian Medical University Ethic Committee ruled that no formal ethics approval
273 was required in this particular case.

274 **Consent for publication**

275 Not applicable

276 **Availability of data and materials**

277 The datasets used and/or analysed during the current study are available from the
278 corresponding author on reasonable request.

279 **Competing interests**

280 The authors declare that they have no conflict of interest.

281 **Funding:**

282 This work was supported during manuscript writing and language editing by
283 Science and Technology Planning Project of Quanzhou Science and Technology

284 Bureau grant number [2018N001S] and Fujian Provincial Health and Family planning
285 Commission Research Talent training project grant number [2018-1-60].

286 **Authors' contributions**

287 RL drafted the manuscript, conducted IMRT plans and collected the data. YW
288 conceived and designed this study, contoured the regions of interest, and revised the
289 manuscript. GY conducted IMRT plans and performed the analysis. YH selected the
290 patient collective and helped to conduct IMRT plans. All authors read and approved
291 the final manuscript.

292 **Acknowledgements**

293 Not applicable

294 **Abbreviation**

295 BC: breast cancer; BCS: breast-conservation surgery; IMRT: Intensity Modulated
296 Radiation Therapy; OAR: organs-at-risk; CT: Computed tomography; ICRU:
297 international Commission on Radiation Units and Measurements; CSTRO: China
298 Society for Radiation Oncology; DMLC: Dynamic Multi-Leaf Collimator; LAD: left
299 anterior descending artery; LV: left ventricle; CTV: clinical target volumes; PTV:
300 planning target volumes; ROI: Region of Interest; HI: homogeneity index; CI:
301 conformity index; RIST: Radiation-induced skin toxicity;

302

303 **Reference**

304 1. Meara SJ, Langmack KA. An investigation into the use of carbon fibre for megavoltage

305 radiotherapy applications. *Physics in medicine and biology*. 1998;43(5):1359-66.

306 2. Sheykhoo A, Abdollahi S, Hadizadeh Yazdi MH, Ghorbani M, Mohammadi M. Effects of
307 Siemens TT-D carbon fiber table top on beam attenuation, and build up region of 6 MV photon
308 beam. *Reports of practical oncology and radiotherapy : journal of Greatpoland Cancer Center*
309 *in Poznan and Polish Society of Radiation Oncology*. 2017;22(1):19-28.

310 3. Vieira SC, Kaatee RS, Dirkx ML, Heijmen BJ. Two-dimensional measurement of photon
311 beam attenuation by the treatment couch and immobilization devices using an electronic portal
312 imaging device. *Med Phys*. 2003;30(11):2981-7.

313 4. De Puyseleer A, De Neve W, De Wagter C. A patient immobilization device for prone
314 breast radiotherapy: Dosimetric effects and inclusion in the treatment planning system. *Phys*
315 *Med*. 2016;32(6):758-66.

316 5. Lee KW, Wu JK, Jeng SC, Hsueh Liu YW, Cheng JC. Skin dose impact from vacuum
317 immobilization device and carbon fiber couch in intensity modulated radiation therapy for
318 prostate cancer. *Med Dosim*. 2009;34(3):228-32.

319 6. Munjal RK, Negi PS, Babu AG, Sinha SN, Anand AK, Kataria T. Impact of 6MV photon
320 beam attenuation by carbon fiber couch and immobilization devices in IMRT planning and
321 dose delivery. *Journal of medical physics*. 2006;31(2):67-71.

322 7. Mulliez T, Gulyban A, Vercauteren T, van Greveling A, Speleers B, De Neve W, et al.
323 Setup accuracy for prone and supine whole breast irradiation. *Strahlentherapie und Onkologie*
324 *: Organ der Deutschen Rontgengesellschaft [et al]*. 2016;192(4):254-9.

325 8. Yu T, Xu M, Sun T, Shao Q, Zhang Y, Liu X, et al. External-beam partial breast irradiation
326 in a supine versus prone position after breast-conserving surgery for Chinese breast cancer

327 patients. Scientific reports. 2018;8(1):15354.

328 9. Harsolia A, Kestin L, Grills I, Wallace M, Jolly S, Jones C, et al. Intensity-modulated
329 radiotherapy results in significant decrease in clinical toxicities compared with conventional
330 wedge-based breast radiotherapy. Int J Radiat Oncol Biol Phys. 2007;68(5):1375-80.

331 10. Kestin LL, Sharpe MB, Frazier RC, Vicini FA, Yan D, Matter RC, et al. Intensity
332 modulation to improve dose uniformity with tangential breast radiotherapy: initial clinical
333 experience. Int J Radiat Oncol Biol Phys. 2000;48(5):1559-68.

334 11. Vicini FA, Sharpe M, Kestin L, Martinez A, Mitchell CK, Wallace MF, et al. Optimizing
335 breast cancer treatment efficacy with intensity-modulated radiotherapy. Int J Radiat Oncol Biol
336 Phys. 2002;54(5):1336-44.

337 12. Roberts KB, Soulos PR, Herrin J, Yu JB, Long JB, Dostaler E, et al. The adoption of new
338 adjuvant radiation therapy modalities among Medicare beneficiaries with breast cancer:
339 clinical correlates and cost implications. Int J Radiat Oncol Biol Phys. 2013;85(5):1186-92.

340 13. Wang EH, Mougalian SS, Soulos PR, Smith BD, Haffty BG, Gross CP, et al. Adoption of
341 intensity modulated radiation therapy for early-stage breast cancer from 2004 through 2011.
342 Int J Radiat Oncol Biol Phys. 2015;91(2):303-11.

343 14. Olson A, Phillips K, Eng T, Lenards N, Hunzeker A, Lewis D, et al. Assessing dose
344 variance from immobilization devices in VMAT head and neck treatment planning: A
345 retrospective case study analysis. Med Dosim. 2018;43(1):39-45.

346 15. Hodapp N. [The ICRU Report 83: prescribing, recording and reporting photon-beam
347 intensity-modulated radiation therapy (IMRT)]. Strahlentherapie und Onkologie : Organ der
348 Deutschen Röntgengesellschaft [et al]. 2012;188(1):97-9.

- 349 16. Zhaozhi Y, Jin M, Jinli M, Xin M, Xingxing C, Xiaomeng Z, et al. Early stage breast cancer
350 postoperative target volume contouring. CHINA ONCOLOGY. 2019;29(09):753-60.
- 351 17. Bellon JR, Wong JS, MacDonald SM, Editors AYH. Radiation Therapy Techniques and
352 Treatment Planning for Breast cancer. Dumane V, Kuo L, Hong L, Ho AY, editors. Springer
353 Nature: Springer International Publishing AG Switzerland; 2016. 20 p.
- 354 18. Oliver M, Chen J, Wong E, Van Dyk J, Perera F. A treatment planning study comparing
355 whole breast radiation therapy against conformal, IMRT and tomotherapy for accelerated
356 partial breast irradiation. Radiother Oncol. 2007;82(3):317-23.
- 357 19. Nakamura JL, Verhey LJ, Smith V, Petti PL, Lamborn KR, Larson DA, et al. Dose
358 conformity of gamma knife radiosurgery and risk factors for complications. Int J Radiat Oncol
359 Biol Phys. 2001;51(5):1313-9.
- 360 20. Olch AJ, Gerig L, Li H, Mihaylov I, Morgan A. Dosimetric effects caused by couch tops
361 and immobilization devices: Report of AAPM Task Group 176. Medical Physics.
362 2014;41(6Part1).
- 363 21. Chen L, Peng YL, Gu SY, Shen H, Zhang DD, Sun WZ, et al. Dosimetric Effects of Head
364 and Neck Immobilization Devices on Multi-field Intensity Modulated Radiation Therapy for
365 Nasopharyngeal Carcinoma. J Cancer. 2018;9(14):2443-50.
- 366 22. McCormack S, Diffey J, Morgan A. The effect of gantry angle on megavoltage photon
367 beam attenuation by a carbon fiber couch insert. Medical Physics. 2005;32(2):483-7.
- 368 23. Pastore F, Conson M, D'Avino V, Palma G, Liuzzi R, Solla R, et al. Dose-surface analysis
369 for prediction of severe acute radio-induced skin toxicity in breast cancer patients. Acta Oncol.
370 2016;55(4):466-73.

371 24. Lee TF, Sung KC, Chao PJ, Huang YJ, Lan JH, Wu HY, et al. Relationships among
372 patient characteristics, irradiation treatment planning parameters, and treatment toxicity of
373 acute radiation dermatitis after breast hybrid intensity modulation radiation therapy. PloS one.
374 2018;13(7):e0200192.

375

376

377

378

379

380

381

382

383 **Figure legend**

384 **Fig. 1**, Display of the immobilization devices in the axial (a) and sagittal view (b).

385 The orange portion is couch, the purple portion is the breast-board, the green portion
386 is the chest fixation mask of thermoplastic, the skin contour is displayed in yellow and
387 the PTV is red.

388 **Fig. 2**, (a) Dose difference distribution of Cross-sectional plane for a typical patient
389 with left-side breast cancer after BCS. Dose difference was calculated by subtracting
390 Plan- from Plan+. (b) DVH results of Plan- and Plan+ for one typical patient with
391 left-side breast cancer after BCS. The solid lines represent the results of Plan-
392 (calculated without immobilization devices), and the dotted lines represent the results

393 of Plan+ (calculated with the whole immobilization devices included in the external
394 body structure).

395 **Fig. 3,** The \bar{D} % of skin dosimetries for breast cancer after radical mastectomy (a) and
396 after BCS (b). Error bars reflect the standard error of the mean (σ/\sqrt{n}). The lines are
397 drawn only to guide the eye.

398 **Table 1.** Dosimetric parameters of PTV and OARs of left breast cancer after radical
399 mastectomy.

400 **Table 2.** Dosimetric parameters of PTV and OARs of right breast cancer after radical
401 mastectomy.

402 **Table 3.** Dosimetric parameters of PTV and OARs of left breast cancer after
403 breast-conserving surgery.

404 **Table 4.** Dosimetric parameters of PTV and OARs of right breast cancer after
405 breast-conserving surgery.

406

Figures

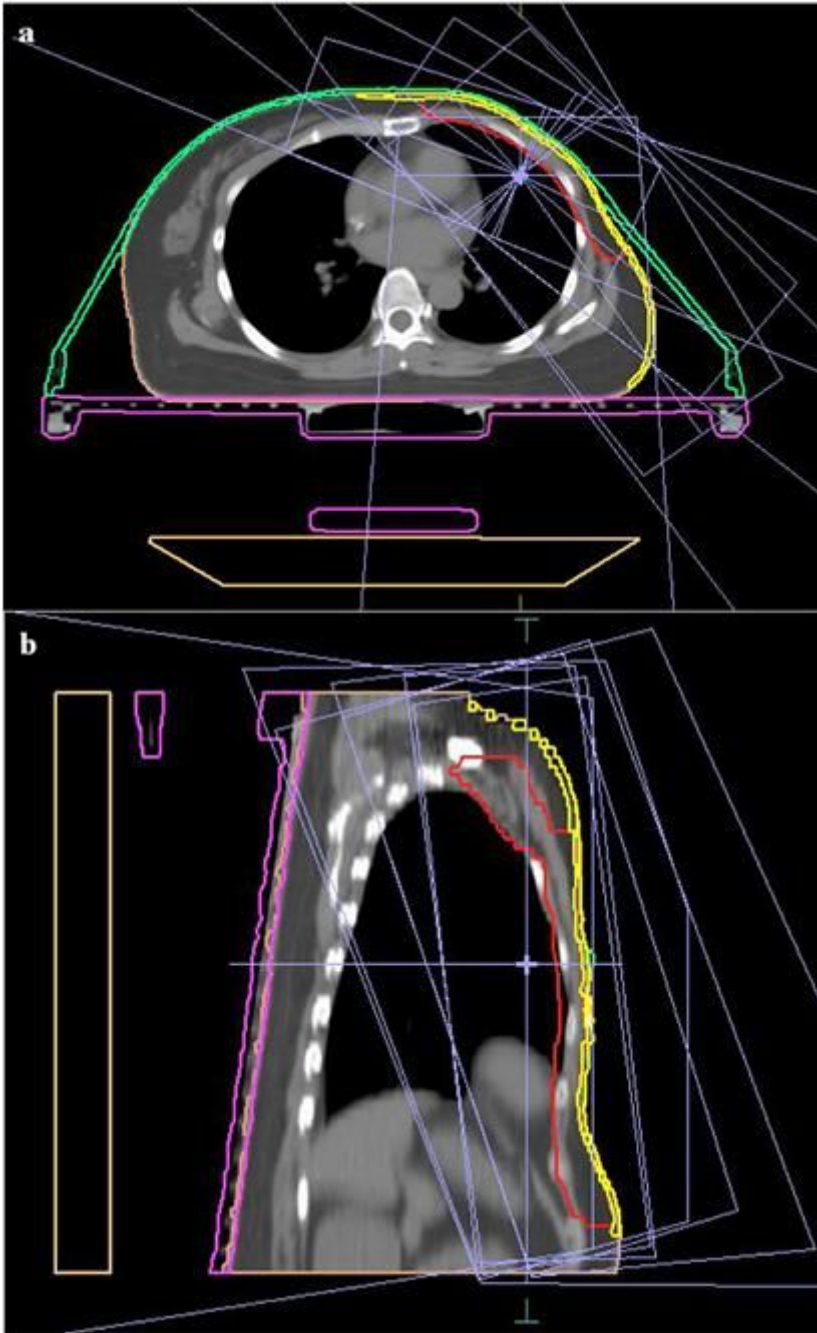


Figure 1

Display of the immobilization devices in the axial (a) and sagittal view (b). The orange portion is couch, the purple portion is the breast-board, the green portion is the chest fixation mask of thermoplastic, the skin contour is displayed in yellow and the PTV is red.

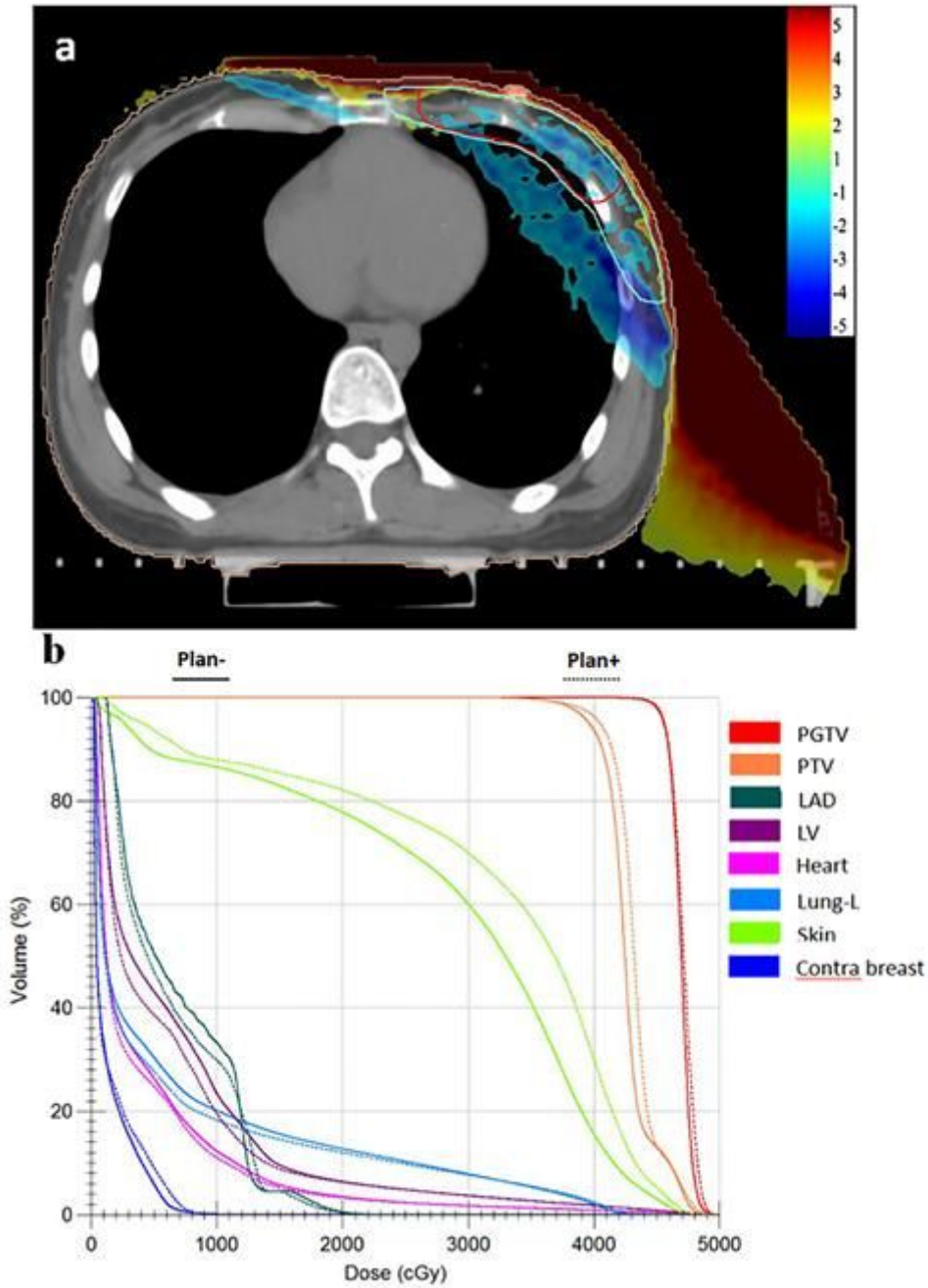


Figure 2

(a) Dose difference distribution of Cross-sectional plane for a typical patient with left-side breast cancer after BCS. Dose difference was calculated by subtracting Plan- from Plan+. (b) DVH results of Plan- and Plan+ for one typical patient with left-side breast cancer after BCS. The solid lines represent the results of Plan- (calculated without immobilization devices), and the dotted lines represent the results of Plan+ (calculated with the whole immobilization devices included in the external body structure).

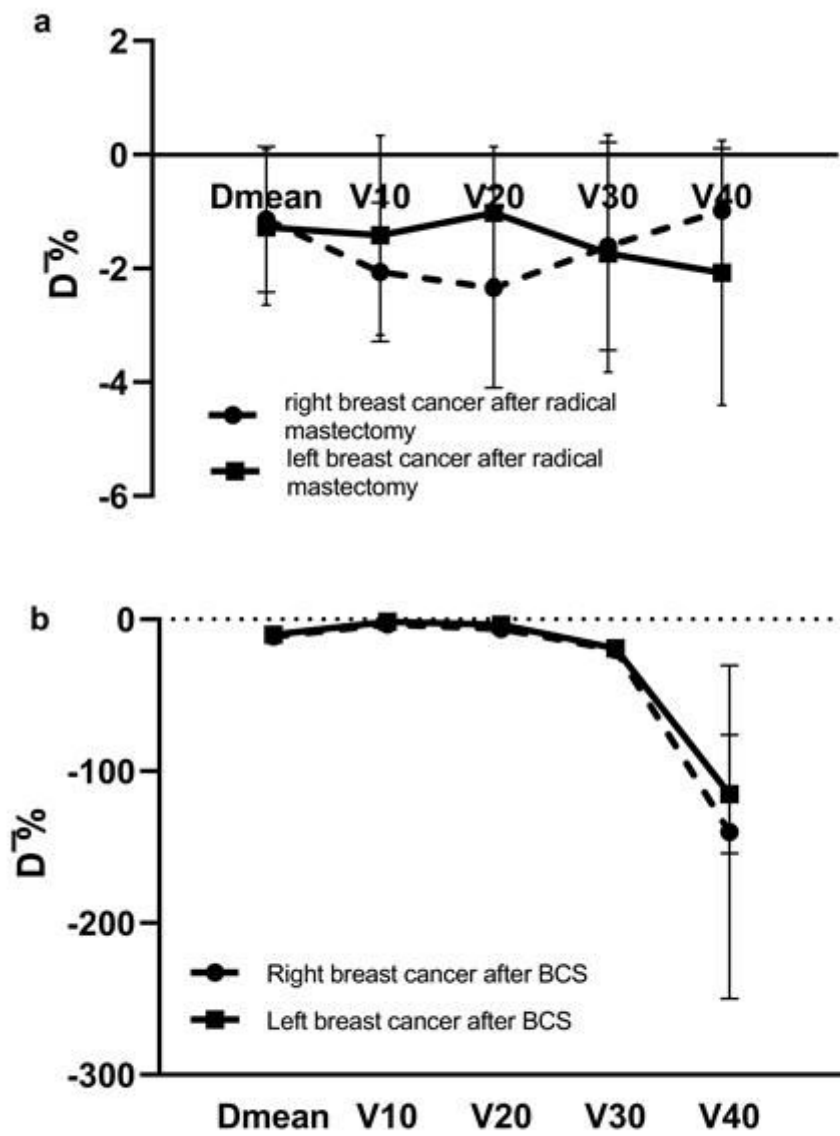


Figure 3

The D% of skin dosimetries for breast cancer after radical mastectomy (a) and after BCS (b). Error bars reflect the standard error of the mean (r/n). The lines are drawn only to guide the eye.