Evaluation of deep learning-based autosegmentation in breast cancer radiotherapy

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Abstract
Purpose: To study the performance of a proposed deep learning-based autocontouring system in delineating organs at risk (OARs) in breast radiotherapy with a group of experts.

Methods: Eleven experts from two institutions delineated nine OARs in 10 cases of adjuvant radiotherapy after breast-conserving surgery. Autocontours were then provided to the experts for correction. Overall, 110 manual contours, 110 corrected autocontours, and 10 autocontours of each type of OAR were analyzed. The Dice similarity coefficient (DSC) and Hausdorff distance (HD) were used to compare the degree of agreement between the best manual contour (chosen by an independent expert committee) and each autocontour, corrected autocontour, and manual contour. Higher DSCs and lower HDs indicated a better geometric overlap. The amount of time reduction using the autocontouring system was examined. User satisfaction was evaluated using a survey.

Results: Manual contours, corrected autocontours, and autocontours had a similar accuracy in the average DSC value (0.88 vs. 0.90 vs. 0.90). The accuracy of autocontours ranked the second place, based on DSCs, and the first place, based on HDs among the manual contours. Interphysician variations among the experts were reduced in corrected autocontours, compared to variations in manual contours (DSC: 0.89–0.90 vs. 0.87–0.90; HD: 4.3–5.8 mm vs. 5.3–7.6 mm). Among the manual delineations, the breast contours had the largest variations, which improved most significantly with the autocontouring system. The total mean times for nine OARs were 37 min for manual contours and 6 min for corrected autocontours. The results of the survey revealed good user satisfaction.

Conclusions: The autocontouring system had a similar performance in OARs as that of the experts’ manual contouring. This system can be valuable in improving the quality of breast radiotherapy and reducing interphysician variability in clinical practice.

Keywords: Autocontouring, Breast, Organs at risk, Radiotherapy

Background
In breast cancer radiotherapy, techniques have evolved from two-dimensional (2D) radiotherapy planning to conformal radiotherapy planning and intensity-modulated radiation therapy [1]. Delineating and sparing organs at risk (OARs) has accordingly received attention recently in breast radiotherapy. In addition, three-dimensional (3D) computed tomography (CT)-based planning, which allows for the accurate assessment of each OAR receiving a radiation dose, has been increasingly used in modern radiotherapy. However, the delineation of OARs is a time-consuming and labor-intensive process and is prone to observer subjectivity, which results in interphysician variations.
With recent advances in big data collection and computing power, deep learning algorithms and procedures have increasingly been used in many different fields [2]. Medical image semantic segmentation, which relies on deep convolutional neural networks, has been extensively studied [3]. Unlike other image segmentation used in surgical and radiologic fields, normal tissue contouring in radiation oncology, known as “OAR delineation,” has been defined and standardized through expert consensus with regard to better quantification of dose-volume histogram–toxicity relationships [4]. Men et al. [5] previously developed deep learning-based target volume for breast radiotherapy, while Feng et al. [6] developed deep learning-based segmentation of OARs for thoracic radiotherapy. Our group also previously demonstrated the potential of deep learning-based autosegmentation of target volumes and OARs in breast cancer radiotherapy [7, 8]. A training set for a proposed deep learning-based autocontouring system (ACS) is generally generated by a single expert or a small group of experts [9]. Therefore, generalization is often discussed as an issue of external validity.

For application of the ACS in real-world clinical practice, its validation with experts from diverse clinical backgrounds, in terms of accuracy, time saving, and user satisfaction, would be necessary. However, our previous studies did not focus on the generalizability or real-world use [7, 8]. In this study, we evaluated the performance of a proposed ACS in delineating OARs for breast radiotherapy with a group of experts from multiple institutions.

Methods

ACS development

Training methods for deep neural networks have been described previously [7]. A home-made ACS was developed. Briefly, a single expert contoured the target volumes and OARs of 111 breast cancer patients who had received adjuvant radiotherapy after breast-conserving surgery. A three-dimensional (3D) U-Net-like convolutional neural network (CNN) was used, which was based on the U-Net structure [10], and combined with 3D version of EfficientNet-B0 as the backbone. Among the 111 cases, 92 were used as the training dataset, and 19 were used as the test dataset. Quantitative tests included the Dice similarity coefficient (DSC) and 95% Hausdorff distance (HD) and revealed an acceptable correlation between the autosegmented and manual contours. Qualitative tests included the scoring, after other panels reviewed the autocontours, and revealed acceptable results.

Study design

There were 11 experts with a median of 7 years (range 2–21 years) of experience in breast cancer radiotherapy who volunteered to participate in this study. The experts were attending physicians (n = 2), clinical fellows (n = 6), residents (n = 2), and a dosimetrist (n = 1) from two institutions (Yonsei Cancer Center [Seoul, South Korea] and Asan Medical Center [Seoul, South Korea]). First, the 11 experts manually delineated the OARs (thyroid, right/left lung, spinal cord, esophagus, heart, liver, and right/left breast) in breast cancer radiotherapy on simulation CT scans of 10 women planning to undergo radiotherapy for breast cancer (i.e., manual contours). Second, an ACS was used on the same simulation CT scans, and these autocontours were provided to the experts. The experts were asked to correct the autocontours, as needed (i.e., corrected autocontours). Before contouring, CT scans were de-identified and the patients’ clinical information was blinded. The clinical treatment contours used for the patient’s radiotherapy delivery were removed to avoid bias during contouring. The experts were asked to record a video during contouring for each CT scan by using screen-recording software (oCam; OHSOFT, South Korea).

The best manual contours for each simulation CT were then selected as the ground truth, after a blind review of contour images of all CT slices by an independent third-party committee. The committee comprised five attending physicians in the radiation oncology department who were breast cancer authorities, and no member was part of the delineation group. Each member scored the performance of each contour, and ground truth was determined by the highest sum of scores. The second-best manual contours were determined by the second-highest sum of scores. This blind review of contours was conducted online by using a questionnaire platform by Google (Menlo Park, CA). By using these ground truths, accuracy was compared between the manual autocontour, corrected autocontour, and autocontour groups.

Endpoints

Endpoints were determined based on three aspects: (1) accuracy of OARs volumes and interphysician variability, (2) time-saving effect, and (3) user satisfaction. To assess accuracy of OARs volumes and interphysician variability, the DSC and 3D HD were used; a higher DSC and a lower HD indicated better geometric overlap. DSC was defined as $D(A, B) = 2|A \cap B|/(|A| + |B|)$ and describes the relative overlap of segmentation volumes A and B. The DSC values range from 0 to 1 with a score of 0 indicating no overlap and 1 indicating perfect overlap. In addition, the HD was used to assess the amount of gross error between
To determine the accuracy of OARs volumes, DSCs and HDs were compared between manual contours, corrected autocontours, and autocontours using the paired t-test. For group-wise comparisons, *P*-values were corrected with Bonferroni's method to counter the problem of multiple comparison. Values of *P* < 0.05 were considered statistically significant. Statistical calculations were conducted using SPSS software (version 25; IBM, Armonk, NY) and GraphPad Prism Version 8 (GraphPad Software, San Diego, CA).

### Results

#### Accuracy

We collected 110 manual contours, 110 corrected autocontours, and 10 autocontours of each type of OAR. When these contours were compared to the consensus ground truth contours, 100 DSCs and 100 HDs (i.e., pairs of the ground truth contour and each contour) were created for each type of OAR for the manual contours and corrected autocontours, and 10 DSCs and 10 HDs were created for the autocontours.

Table 1 and Fig. 1 show the mean DSCs and HDs of the manual contours, corrected autocontours, and autocontours for each OAR. In general, manual contours, corrected autocontours, and autocontours had a

### Table 1  Summary of DSC and HD

| Contour  | (1) Manual (n = 100) | (2) Corrected auto (n = 100) | (3) Auto (n = 10) | P-value* |
|---------|---------------------|-----------------------------|-----------------|----------|
| **DSC (mean ± SE)** |                     |                             |                 |          |
| Thyroid  | 0.80 ± 0.01         | 0.80 ± 0.01                 | 0.79 ± 0.02     | .953     |
| Lung_right | 0.98 ± 0.00       | 0.97 ± 0.00                 | 0.97 ± 0.00     | < .001  |
| Lung_left | 0.97 ± 0.00        | 0.96 ± 0.00                 | 0.96 ± 0.01     | < .001   |
| Breast_right | 0.81 ± 0.00   | 0.90 ± 0.00                 | 0.91 ± 0.00     | < .001   |
| Breast_left | 0.83 ± 0.00      | 0.90 ± 0.00                 | 0.90 ± 0.01     | < .001   |
| Spinal cord | 0.82 ± 0.01     | 0.85 ± 0.00                 | 0.85 ± 0.01     | .001     |
| Esophagus | 0.84 ± 0.00        | 0.83 ± 0.00                 | 0.82 ± 0.01     | .001     |
| Heart    | 0.92 ± 0.00        | 0.94 ± 0.00                 | 0.95 ± 0.00     | < .001   |
| Liver    | 0.94 ± 0.00        | 0.94 ± 0.00                 | 0.93 ± 0.01     | < .001   |
| **HD (mean ± SE)** |                     |                             |                 |          |
| Thyroid  | 3.82 ± 0.20        | 4.12 ± 0.26                 | 4.28 ± 0.83     | .33      |
| Lung_right | 2.37 ± 0.15       | 2.46 ± 0.09                 | 2.42 ± 0.29     | > .999   |
| Lung_left | 3.61 ± 0.35        | 2.93 ± 0.18                 | 2.99 ± 0.67     | 0.077    |
| Breast_right | 12.44 ± 0.80 | 8.06 ± 0.35                 | 7.54 ± 0.65     | < .001   |
| Breast_left | 10.85 ± 0.58    | 8.14 ± 0.33                 | 8.15 ± 1.05     | < .001   |
| Spinal cord | 3.95 ± 0.54     | 2.20 ± 0.04                 | 2.21 ± 0.11     | 0.005    |
| Esophagus | 3.95 ± 0.36        | 3.16 ± 0.06                 | 3.46 ± 0.21     | 0.1      |
| Heart    | 8.26 ± 0.55        | 5.42 ± 0.29                 | 4.73 ± 0.33     | < .001   |
| Liver    | 6.92 ± 0.64        | 5.95 ± 0.32                 | 9.74 ± 3.16     | 0.335    |

Abbreviations: DSC, Dice similarity coefficient; HD, Hausdorff distance; SE, standard error

* *P*-values were calculated using the paired t-test with Bonferroni correction
similar accuracy from the average DSCs (0.88 vs. 0.90 vs. 0.90). However, for breast contours, the DSCs were significantly higher in the corrected autocontours and autocontours than in the manual contours. The corrected autocontours of the breast had better DSCs than the manual contours by 0.09 (right breast) and 0.07 (left breast). In contrast, the absolute difference in the DSCs between the corrected autocontours and manual contours for other OARs was relatively small. For example, the absolute difference in DSCs between the two groups was 0.01 for the lungs, < 0.01 for the thyroid, 0.03 for the spinal cord, 0.01 for the esophagus, 0.02 for the heart, and < 0.01 for the liver.

The HDs of breast and heart contours were significantly lower in the corrected autocontours and autocontours than in the manual contours. The HD of liver contours were significantly higher in the autocontours than in the corrected autocontours. The results of the sensitivity analyses were consistent with the original analyses for all OARs, excluding the thyroid and lungs (Additional file 1: Table S1 and Additional file 1: Figure S1).

To evaluate the performance of autocontouring alone, the average DSCs and HDs of all nine OARs were compared between manual contours and autocontours. In the manual contours, the average values of the DSCs of all OARs ranged from 0.870 to 0.903 (median, 0.881), depending on the expert, and the average values of HDs of all OARs ranged from 5.327 mm to 7.636 mm (median, 6.431 mm). Based on these DSCs, autocontours ranked second with a value of 0.896, after the manual contour’s value of 0.903. Based on these HDs, autocontours ranked first place with a value of 5.142 mm, followed by manual contours with a value 5.327 mm (Table 2).

Interphysician variability

The interphysician variations observed in the experts’ manual contours were reduced in the corrected autocontours. The mean DSCs of all OARs ranged from 0.87 to 0.90, based on the individuals’ manual contours, although the range was reduced to 0.89–0.90 in the individuals’ corrected autocontours. The mean HDs of all OARs ranged from 5.3 mm to 7.6 mm, based on the individuals’ manual contours, although the range was reduced to 4.3–5.8 mm in the individuals’ corrected autocontours. Figure 2 shows the mean DSCs based on OARs. This figure shows that DSCs were more homogeneous in the corrected autocontours than in the
manual contours, indicating reduced interphysician variability. The sensitivity analysis in Additional file 1: Figure S2 reveals results consistent to those of the original analyses.

Examples of manual and corrected autocontours of the breast and heart are shown in Fig. 3. Of note, the interphysician variability of manual breast contours mostly occurred in the lateral and anterior borders of the breast, whereas this variability rarely occurred in the corrected autocontours. For heart contours, interphysician variability of the manual contours mostly occurred in the superior borders, whereas this variability rarely occurred in the corrected autocontours.

Time-saving effect
The mean (± standard error) contouring time for the nine OARs of each patient was 37.4 (± 5.9) min for the manual contours and 6.4 (± 1.4) min for corrected autocontours, which indicated a time reduction of 84% with the ACS (Fig. 4A and Additional file 1: Table S2). The process of obtaining autocontours was fully automated and took < 10 min, depending on the computer performance. When the mean time was measured, based on each OAR, breast and liver contouring was the longest step among the manual contours. The time was prominently reduced in the corrected autocontours [right breast: from 5.9 (± 1.2) min to 0.5 (± 0.3) min; left breast: from 6.3 (± 1.2) min to 0.6 (± 0.2) min; liver: from 9.0 (± 1.5) min to 1.5
User satisfaction

The mean (± standard error) scores of questions regarding user satisfaction were as follows: question 1, 7.5 (± 0.3); question 2, 8.8 (± 0.3); and question 3, 9.2 (± 0.2).

Discussion

Deep learning-based autosegmentation has been widely investigated in head and neck, thoracic, and genitourinary malignancies, although there is relatively less data regarding the use of deep learning-based autosegmentation in breast radiotherapy planning, for which conventional field-based and 2D radiotherapy techniques are commonplace. In this context, we previously developed a deep learning-based ACS for breast radiotherapy planning and reported its feasibility [7]. The clinical target volumes for the breasts and OARs were manually contoured in simulation CT scans and trained with a 3D U-Net-like CNN.

By using our deep learning-based ACS, we compared the performance between manual contours, corrected autocontours, and autocontours with multiple experts from two institutions. We showed that autocontours and corrected autocontours were closer to the ground truth than were manual contours. Furthermore, the accuracy of autocontours for breasts and OARs was as good as that of other deep learning-based ACSs developed by Men et al. [5] and Feng et al. [6]. The interphysician variation in manual contours was greatly reduced with the ACS. Moreover, the time spent contouring was substantially reduced with the ACS. Satisfaction was good among participants using the ACS.

Before the era of deep learning-based autosegmentation, atlas-based autosegmentation was actively studied in breast cancer and in other cancer sites. In a previous study [8], we compared the performance of deep learning-based autosegmentation with that of atlas-based autosegmentation in breast OARs and clinical target volumes. The proposed deep learning-based autosegmentation showed more consistent results and outperformed atlas-based autosegmentation in most structures. The next clinically relevant issue would be to address how the performance of the proposed deep-learning algorithm, which was trained by a single expert, would compare with that of a group of experts.

The novelty of the current study is that experts’ manual contours and autocontours were compared with the ground truth, which was determined by the other third-party experts. The average DSC and HD of autocontours ranked second and first place, respectively, in relation to the experts’ manual contours. This finding indicated that the autocontours have, at least, a similar performance as that of the experts’ manual contours. The good performance of autocontours may be because of the good quality of the training dataset for deep learning systems attached to the contouring guidelines [4]. In addition, possible human errors in manual contouring may be caused by fatigue from repetitive work. Another strength of this study is the integration of experts with diverse clinical backgrounds and participation of different institutions.

We observed substantial interphysician variability between the experts’ manual contours. Substantial variability in the manual contouring of the targets and OARs between institutions and observers was demonstrated in a Radiation Therapy Oncology Group (RTOG)
multi-institutional and multiobserver study [12]. Such interphysician variability is an obstacle in accurately assessing the efficacy of radiotherapy and risk of long-term adverse effects. Incidental radiation exposure to the heart during breast radiation therapy increases the risk of heart disease with regard to the dose–response relationship between heart radiation dose and an acute coronary event [13, 14]. Moreover, radiation-related hypothyroidism [15], radiation pneumonitis [16], and secondary contralateral breast cancer [17] have been reported in patients with breast cancer. In addition, in clinical trials including radiotherapy, standardization of treatment is problematic because of the variability in delineating the target and OARs [18]. In the RTOG 0617 trial [19], a radiation dose-escalation trial of non-small cell lung cancer, an analysis using deep-learning segmented hearts revealed that the actual heart doses were higher than originally reported owing to inconsistent and insufficient manual heart segmentation. Our results demonstrated that the ACS could solve this issue. For example, the lateral border of the breast had the largest variation among the experts’ manual contours in this study. The most widely used RTOG guidelines [20] and ESTRO guidelines [21] define the lateral border of the breast as a clinically palpable breast or lateral breast fold; therefore, clearly defining it on a CT image is difficult. The ACS can aid in standardizing delineation when the definition of the boundary of an organ is ambiguous.

For the breast and heart contours, the autocontours and corrected-autocontours had a significantly better accuracy than did the manual contours. By contrast, the HDs of liver contours were high in the autocontours and reduced in the corrected autocontours to a value similar to that of manual contours, suggesting that manual adjustment was necessary. Therefore, the detailed performance of ACS appears to vary depending on the OAR.

The manual adjustment of autocontouring had an average time reduction of 84%, compared with manual contouring. This was most remarkable in breast and liver contouring, which required the most time. When adjusting the autocontour, the average time taken for each organ was <1 min, indicating that only minimal or no correction was needed. In addition, the participants responded that ACS helped to shorten the time spent contouring and that they would like to use it in the future. According to Shanafelt et al. [22], symptoms of burnout have been reported in >50% of practicing physicians, and this affliction is largely driven by work-related stressors [23]. Therefore, efforts are needed to reduce the workload of physicians. The ACS can be used for this.

This study has a limitation. It is difficult to determine the ultimate ground truth. Although the ground truth was determined by a separate qualified group of attending physicians, other experts may not agree with the ground truth. Thus, to clarify the results, we conducted the same analysis by using the second-best contours instead of the ground truth contours for the sensitivity analysis. The results were generally consistent with the original analyses for all OARs, excluding the thyroid and lungs.

Conclusions
The ACS can overcome several weaknesses of manual contouring, such as labor intensity, time consumption, and interphysician variation. To expand the frame of this study, we are conducting a multicenter study in the Korean Radiation Oncology Group to examine the effectiveness of the ACS in the breast target. In the future, delineating OARs and accurately assessing the toxicity risk by the irradiated dose of each organ will become more important in breast cancer radiotherapy as treatments become more sophisticated. Adopting the ACS in breast cancer radiotherapy could be helpful in this regard.

Abbreviations
2D: Two-dimensional; 3D: Three-dimensional; ACS: Autocontouring system; CT: Computed tomography; DSC: Dice similarity coefficient; HD: Hausdorff distance; OAR: Organ at risk; RTOG: Radiation Therapy Oncology Group.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s13014-021-01923-1.

Acknowledgements
We thank Minji Koh, Young Seob Shin, and Jesang Yu of the Department of Radiation Oncology at Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea, and Jina Kim, Taehyung Kim, Jason Joon Bok Lee, Jin Young Moon, and Ryeong Hwang Park of the Department of Radiation Oncology at Yonsei University College of Medicine, Seoul, Korea, for participating in contouring.

Authors’ contributions
Study conception and design: H.K.B., J.S.C., JJ, YBK; provision of study materials: J.S.C, J.J, C.J., J.S.K., SYC, SL, YBK; collection and assembly of data: H.K.B.; data analysis and interpretation: H.K.B., J.S.C., M.S.C., J.C., JJ; manuscript writing: H.K.B., J.S.C., JJ. All authors read and approved the final manuscript.
Funding
This work was supported by a National Research Foundation of Korea (NRF) grant, funded by the Korea government (MSIT; Grant No. 2019R1C1C009359).

Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate
This study was approved by our Institutional Review Board (4-2020-0466). Informed consent was waived.

Consent for publication
All authors have approved the manuscript and agree with submission to Radiation Oncology.

Competing interests
None.

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Received: 30 May 2021 Accepted: 27 September 2021 Published online: 14 October 2021

References
1. Fogliata A, Nicolini G, Alber M, Asell M, Dobler B, El-Haddad M, et al. IMRT for breast: A planning study. Radiother Oncol. 2005;76(3):300–10.
2. Bi WL, Hosny A, Schabath MB, Giger ML, Birkbak NJ, Mehrtash A, et al. Artificial intelligence in cancer imaging: clinical challenges and applications. CA Cancer J Clin. 2019;69(2):127–57.
3. Jiang F, Grigorev A, Rho S, Tian Z, Fu Y, Jifara W, et al. Medical image semantic segmentation based on deep learning. Neural Comput Appl. 2018;29(5):1257–65.
4. Wright JJ, Yom SS, Awan MJ, Dawes S, Fischer-Valuck B, Kudner R, et al. Standardizing normal tissue contouring for radiotherapy treatment planning: an ASTRO consensus paper. Pract Radiat Oncol. 2019;9(2):65–72.
5. Men K, Zhang T, Chen X, Chen B, Tang Y, Wang S, et al. Fully automatic and robust segmentation of the clinical target volume for radiotherapy of breast cancer using big data and deep learning. Phys Med. 2018;50:13–9.
6. Feng X, Qinq K, Tutson JD, Meyer CH, Chen Q. Deep convolutional neural network for segmentation of thoracic organs-at-risk using cropped 3D images. Med Phys. 2019;46(5):2160–80.
7. Chung SY, Chang JS, Choi MS, Chang Y, Choi BS, Chun J, et al. Clinical feasibility of deep learning-based auto-segmentation of target volumes and organs-at-risk in breast cancer patients after breast-conserving surgery. Radiat Oncol. 2021;16(1):1–10.
8. Choi M5, Choi BS, Chung SY, Kim N, Chun J, Kim YB, et al. Clinical evaluation of atlas- and deep learning-based automatic segmentation of multiple organs and clinical target volumes for breast cancer. Radiother Oncol. 2020;153:139–45.
9. Liu Z, Liu X, Guan H, Zhen H, Sun Y, Chen Q, et al. Development and validation of a deep learning algorithm for auto-delineation of clinical target volume and organs at risk in cervical cancer radiotherapy. Radiother Oncol. 2020;153:172–9.
10. Ronneberger O, Fischer P, Brox T. U-net: Convolutional networks for biomedical image segmentation. In: International conference on medical image computing and computer-assisted intervention. Springer; 2015. p. 234–41.
11. Kumarasiri A, Siddiqui F, Liu C, Yecheili R, Shah M, Pradhan D, et al. Deformable image registration based automatic CT-to-CT contour propagation for head and neck adaptive radiotherapy in the routine clinical setting. Med Phys. 2014;41(12):121712.
12. Li YA, Tai A, Arthur DW, Buchholz TA, Macdonald S, Marks LB, et al. Variability of target and normal structure delineation for breast cancer radiotherapy: an RTOG Multi-Institutional and Multiobserver Study. Int J Radiat Oncol Biol Phys. 2009;73(3):944–51.
13. Chung SY, Oh J, Chang JS, Shin J, Kim KH, Chun KH, et al. Risk of cardiac disease in breast cancer patients: impact of patient-specific factors and individual heart dose from three-dimensional radiotherapy planning. Int J Radiat Oncol Biol Phys. 2021. https://doi.org/10.1016/j.ijrobp.2020.12.053.
14. Darby SC, Ewertz M, McGale P, Bennet AM, Blom-Goldman U, Brennum D, et al. Risk of ischemic heart disease in women after radiotherapy for breast cancer. N Engl J Med. 2013;368(1):987–98.
15. Choi SH, Chang JS, Son NH, Hong CS, Byun HK, Hong N, et al. Risk of hypothyroidism in women after radiotherapy for breast cancer. Int J Radiat Oncol Biol Phys. 2019. https://doi.org/10.1016/j.ijrobp.2019.09.047.
16. Choi J, Kim YB, Shin KH, Ahn SJ, Lee HS, Park W, et al. Radiation pneumonitis in association with internal mammary node irradiation in breast cancer patients: an ancillary result from the KROG08–06 study. J Breast Cancer. 2016;19(3):275–82.
17. Zhang Q, Liu J, Ao N, Yu H, Peng Y, Ou L, et al. Secondary cancer risk after radiotherapy for breast cancer with different radiotherapy techniques. Sci Rep. 2020;10(1):1220.
18. Pérez CA, Gardner P, Glasgow GP. Radiotherapy quality assurance in clinical trials. Int J Radiat Oncol Biol Phys. 1984;10(Suppl 1):119–25.
19. Thor M, Apte A, Haq R, Iyer A, LoCastro E, Deasy JO. Using auto-segmentation to reduce contouring and dose inconsistency in clinical trials: the simulated impact on RT0G 0617. Int J Radiat Oncol Biol Phys. 2021;109(5):1619–26.
20. Sun GY, Wang SL, Song YW, Jin J, Wang WH, Liu YP, et al. Radiation-induced lymphopenia predicts poorer prognosis in patients with breast cancer: a post hoc analysis of a randomized controlled trial of postmastectomy hypofractionated radiation therapy. Int J Radiat Oncol Biol Phys. 2020;108(1):277–85.
21. Offersen BV, Boersma LJ, Kirkove C, Hol S, Aznar MC, Sola AB, et al. ESTRO consensus guideline on target volume delineation for elective radiotherapy of early stage breast cancer, version 1.1. Radiother Oncol. 2020;153:172–9.
22. Shanafelt TD, Hasan O, Dyrbye LN, Sinsky C, Satele D, Sloan J, et al. Association of clinical specialty with symptoms of burnout and career choice regret among US resident physicians. JAMA. 2016;315(1):114–30.

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