



# **Special Article**

# Standardizing Normal Tissue Contouring for Radiation Therapy Treatment Planning: An ASTRO Consensus Paper



Jean L. Wright MD <sup>a</sup>, Sue S. Yom MD, PhD, MAS <sup>b</sup>, Musaddiq J. Awan MD <sup>c</sup>, Samantha Dawes CMD <sup>d,\*</sup>, Benjamin Fischer-Valuck MD <sup>e</sup>, Randi Kudner MA <sup>d</sup>, Raymond Mailhot Vega MD, MPH <sup>f</sup>, George Rodrigues MD, PhD <sup>g</sup>

<sup>a</sup>Johns Hopkins University, Baltimore, Maryland; <sup>b</sup>University of California, San Francisco, California; <sup>c</sup>Medical College of Wisconsin, Milwaukee, Wisconsin; <sup>d</sup>American Society for Radiation Oncology, Arlington, Virginia; <sup>e</sup>Emory University, Atlanta, Georgia; <sup>f</sup>University of Florida, Jacksonville, Florida; <sup>g</sup>London Health Sciences Centre, London, ON, Canada

Received 4 September 2018; revised 29 November 2018; accepted 8 December 2018

#### Abstract

**Purpose:** The comprehensive identification and delineation of organs at risk (OARs) are vital to the quality of radiation therapy treatment planning and the safety of treatment delivery. This guidance aims to improve the consistency of ontouring OARs in external beam radiation therapy treatment planning by providing a single standardized resource for information regarding specific OARs to be contoured for each disease site. The guidance is organized in table format as a quality assurance

Note—Earn CME credit by taking a brief online assessment at https://www.astro.org/JournalCME.

Disclaimer and Adherence: ASTRO clinical practice statements present scientific, health, and safety information and may reflect scientific or medical opinion. They are available to ASTRO members and the public for educational and informational purposes only. Commercial use of any content in this paper without the prior written consent of ASTRO is strictly prohibited.

Adherence to this guidance will not ensure successful treatment in every situation and should not be deemed inclusive of all proper methods of care or exclusive of other methods reasonably directed to obtaining the same results. The physician must make the ultimate judgment regarding any specific therapy in light of all circumstances presented by the patient. ASTRO assumes no liability for the information, conclusions, and findings contained in its papers. This guidance cannot be assumed to apply to the use of these interventions performed in the context of clinical trials.

This guidance is based on information available at the time the task force conducted its research and discussions on this topic. There may be new developments that are not reflected in this guidance and that may, over time, be a basis for ASTRO to revisit and update the paper.

Sources of support: This paper was funded by the American Society for Radiation Oncology.

Conflicts of interest: Before initiation of this paper, all members of the Task Force and Review Panel completed disclosure statements. These statements are maintained at ASTRO Headquarters in Arlington, Virginia. ASTRO's Clinical Affairs and Quality Council reviewed these disclosures and determined they do not present a conflict with respect to the Task Force members' work on this consensus paper. Jean Wright: ASTRO (honoraria), International Journal of Radiation Oncology Biology Physics (Associate Editor); Sue S. Yom: ASTRO (honoraria), Genentech, Merck, Bristol-Myers Squibb (research grants), UpToDate, Springer (royalty), American Radium Society (Executive Committee Treasurer, Appropriate Use Criteria Committee Chair), International Journal of Radiation Oncology Biology Physics (Deputy Editor); Raymond Mailhot Vega: Varian (travel expenses) and IBA (travel expenses), George Rodrigues: Demos Publishing Company (royalty), George Rodrigues Medicine Professional Corporation (President, stock).

Supplementary Materials: Tables E1 and E2. Table E1 contains two designations for anatomic sites in the EBRT setting: recommended (structures that are recommended for adult definitive cases and may inform palliative cases) and consider (structures that may be considered depending on the specific clinical scenario). Table E2 is a reference guide for accessing site-specific sources to assist with the delineation of the structures from Table E1.

\* Corresponding author. American Society for Radiation Oncology, 251 18th Street South, 8th Floor, Arlington, VA 22202. E-mail address: Samantha.Dawes@astro.org (S. Dawes). tool for practices and a training resource for residents and other radiation oncology students (see supplementary materials).

**Methods and Materials:** The Task Force formulated recommendations based on clinical practice and consensus. The draft manuscript was peer reviewed by 16 reviewers, the American Society for Radiation Oncology (ASTRO) legal counsel, and ASTRO's Multidisciplinary Quality Assurance Subcommittee and revised accordingly. The recommendations were posted on the ASTRO website for public comment in June 2018 for a 6-week period. The final document was approved by the ASTRO Board of Directors in August 2018.

**Results:** Standardization improves patient safety, efficiency, and accuracy in radiation oncology treatment. This consensus guidance represents an ASTRO quality initiative to provide recommendations for the standardization of normal tissue contouring that is performed during external beam treatment planning for each anatomic treatment site. Table 1 defines 2 sets of structures for anatomic sites: Those that are recommended in all adult definitive cases and may assist with organ selection for palliative cases, and those that should be considered on a case-by-case basis depending on the specific clinical scenario. Table 2 outlines some of the resources available to define the parameters of general OAR tissue delineation.

**Conclusions:** Using this paper in conjunction with resources that define tissue parameters and published dose constraints will enable practices to develop a consistent approach to normal tissue evaluation and dose documentation.

© 2018 American Society for Radiation Oncology. Published by Elsevier Inc. All rights reserved.

#### Introduction

#### **Background**

Comprehensive identification and delineation of organs-at-risk (OARs) are vital to the quality of radiation therapy treatment planning and the safety of treatment delivery. A lack of standardization for normal tissue contouring allows for variations in practice and potentially affects the quality of dosimetric plans.

A major advance was made in the field of radiation oncology over the past decades with increased dose conformality, particularly as the profession moved from 2- to 3- dimensional to intensity modulated techniques. Increased precision in the delivery of radiation has also been influenced by improvements in patient positioning, advances in on-treatment imaging, and motion management techniques. These improvements have yielded 2 major outcomes: improved avoidance of critical structures and an ability to intensify doses. When targeting tumors, there may be adjacent structures for which clinical decision-making can be affected by toxicity risk depending on disease site, patient age, treatment technique, prior radiation, and other variables. Plan optimization and maximal organ sparing ultimately depend on the specification and standardization of OAR definitions, accuracy of OAR delineation, and accounting for organ motion when applicable.

The goal of this guidance is to improve the consistency of contouring OARs in external beam radiation therapy treatment planning by providing a single standardized resource for information regarding specific OARs to be contoured for each disease site.

#### Scope

The Task Force organized the recommendations into a user-friendly table format as a quality assurance tool for practices and a training resource for residents and other radiation oncology students. Table E1 (available online at https://doi.org/10.1016/j.prro.2018.12.003) contains two designations for anatomic sites in the external beam radiation therapy setting: recommended (structures that are recommended for adult definitive cases and may inform palliative cases) and consider (structures that may be considered depending on the specific clinical scenario).

Understanding that practices and patients need the best outcomes given possible resource constraints, the recommended list is focused on the essential organs needed for each primary site in the definitive setting to provide a basic minimum standard of care. The organs listed in the consider category build on the recommended list and represent options relevant to specific clinical situations. These should be considered by the radiation oncology team depending on clinical factors such as anatomic location, treatment intent (palliative vs definitive), applicable pediatric cases, use of stereotactic radiation therapy, or requirements for a clinical trial. In the recommended category, the structures center around those organs with widely recognized dose limits. However, for research or planning purposes, individual practices may wish to include other contours for which dose limits have not yet been clearly established. Contouring normal tissues enables documentation of the dose, even when the recommended dose constraints are not available.

Table E2 (available online at https://doi.org/10.1016/j.prro.2018.12.003) is a reference guide to access site-specific

sources to assist with the delineation of the structures in Table E1. The list is not exhaustive, and other resources may be available in the medical literature and in clinical trial protocols to assist in OAR delineation for specific clinical scenarios.

Dose constraints for specific OARs are beyond the scope of this paper, but constraints are available using established guidelines published by Emami et al,<sup>1</sup> the American Association of Physicists in Medicine Task Group TG-101 report,<sup>2</sup> and the Quantitative Analysis of Normal Tissue Effects in the Clinic guidelines as examples.<sup>3</sup> Cooperative group protocols for clinical trials involving radiation therapy also provide additional for dose constraints.<sup>4</sup>

The naming convention for tissue labels in Table E1 aligns with the *American Association of Physicists in Medicine Task Group 263: Standardizing Nomenclatures in Radiation Oncology*, <sup>5</sup> which standardized structure names.

#### **Indications and Considerations**

#### Standardization

Dose constraints are an integral part of treatment planning and a part of quality measurement in Medicare billing, but there is a lack of standardization in how organs are selected and contoured across practices and providers. According to ASTRO's publication, *Safety Is No Accident*, "[a] review of target delineation and image segmentation prior to planning deserves more standardization."

Multiple studies have shown variation in the delineation of radiation therapy targets and OARs in such disease sites as the central nervous system, <sup>7,8</sup> head and neck, <sup>9,10</sup> thorax, <sup>11-13</sup> and pelvis. <sup>14,15</sup> Differences in normal-tissue contouring can affect dosimetric plan optimization, dose volume histogram (DVH) results, and potentially normal tissue complication probabilities and clinical outcomes. For example, in a test case from a patient with oropharyngeal cancer, significant interclinician variation in contouring OARs was associated with differences that ranged from -289% to 56% for mean dose, and -22% to 35% for maximum dose. <sup>16</sup>

Similar variability in normal structure delineation was observed for breast cancer radiation therapy planning in a Radiation Therapy Oncology Group (NRG Oncology) multi-institutional study, where structure overlaps between radiation oncologists were as low as 10%, and volume variations had standard deviations up to 60%. Establishing standardized normal tissue contouring assists with dose assessment during radiation therapy planning, particularly in the era of advanced radiation therapy techniques.

#### Safety

In 2011, ASTRO partnered with the American Association of Physicists in Medicine to develop the Radiation Oncology Incident Learning System (RO-ILS), the only

medical specialty society-sponsored incident learning system for radiation oncology. A recent analysis of aggregated RO-ILS data indicates that, to date, 29% of all reported events occurred in treatment planning, with a featured theme of incorrect normal-tissue definition leading to the mis-estimation of radiation dose. From 2016 to 2018, the Radiation Oncology Health Care Advisory Council identified >60 RO-ILS events that contained an issue with contours. These events include plans with critical structures not contoured, and normal tissues incorrectly or incompletely delineated. <sup>17</sup>

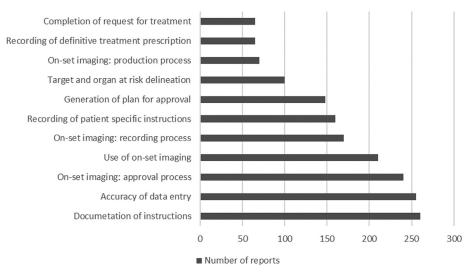
Similarly, reported data of radiation therapy patient events submitted to Public Health England between December 2013 and November 2015 identified that 27% of incidents occurred during the pretreatment planning process. <sup>18</sup> Of those events recorded as near-miss, target and OAR delineation was one of the most frequently reported processes (Fig. 1). This report, along with RO-ILS data, suggests a trend in safety event data relating to treatment planning and promotes the need for improved standardization.

Despite the importance of contouring and potential sources of variation, no recommended quality systems exist to ensure accuracy and consistency in normal anatomy contouring. Standardization may allow better quantification of DVH-toxicity relationships and assist in the prevention of avoidable toxicities to normal structures. Failure to include OARs in the planning process may lead to dose-dumping during the inverse planning process, whereby unnecessary and potentially unsafe radiation doses are delivered to OARs that are not defined as part of the planning constraints. This concept extends to the palliative setting, where doses to OARs are generally considered less critical because of lower radiation doses and shorter patient life expectancy. However, reirradiation and higher-dose stereotactic treatments are increasingly common in the metastatic setting, and if cumulative normal tissue dose constraints are not observed, inadvertent toxicity may occur.

# **Pre-treatment Preparation**

### Physician documentation

Constraining doses to OARs is a fundamental component of the treatment planning process and should be included in the radiation oncologist's planning directive (order) documented before the initiation of treatment planning. The patient-specific written planning directive should be consistently documented in the oncology information system and must provide sufficient information to guide qualified personnel generating a treatment plan. The ASTRO Accreditation Program for Excellence (APEx) Standard 2.2.1<sup>19</sup> requires practices to document defined patient-specific targets and normal tissues along with their respective dose goals and constraints as an essential evaluation component of the program.



**Fig. 1** Public Health England, "Radiotherapy Errors and Near Misses Data Report (December 2013 to November 2015)". <sup>18</sup> Reproduced with permission.

# Roles and responsibilities of treatment planning staff

ASTRO's Safety is No Accident publication highlights the need for all members of the radiation oncology team to maintain proper credentials, skills, and training, and to undergo clinical competency assessments<sup>6</sup> to assure standardization of contouring practices.<sup>20</sup> Ultimately, the radiation oncologist is responsible to review the structures as part of the plan assessment and approval (Fig. 2).

# **Resources**

#### Tools

Contouring guides (i.e., atlases) are published for different anatomic sites to guide practitioners to define clinical target volumes, and should also be used for visualization of tissue parameters. Disease site-specific contouring atlases published by NRG Oncology (commonly known as Radiation Therapy Oncology Group contouring atlases) are developed by consensus among cooperative groups and disease-site committees. Aside from the NRG atlases, other prominent sources exist as published in the medical literature. These tools should be used to support consistency in practice procedures, as teaching resources, and in situations when planning treatment for an uncommon or infrequently treated disease site. Table E2 catalogs some of the available references, and this document seeks to further encourage use of these important tools to promote practice standardization.

#### **Technology**

Modern radiation oncology has experienced rapid changes in software and hardware technology. The ability

|   | Radiation<br>Oncologist | Medical<br>Physicist | Medical<br>Dosimetrist | Radiation<br>Therapist | Nonphysician<br>Providers<br>(NA/PA) | Oncology Nurse |
|---|-------------------------|----------------------|------------------------|------------------------|--------------------------------------|----------------|
| Interdisciplinary coordination of care    | Х                       |                      |                        | Х                      | Х                                    | Х              |
| Patient positioning and image acquisition | Х                       | Х                    | Х                      | Х                      |                                      |                |
| Fusion and registration                   | Х                       | Х                    | Х                      |                        |                                      |                |
| Contouring/segmentation                   | Х                       | Х                    | Х                      |                        |                                      |                |
| Dose-volume constraints                   | Х                       | Х                    | Х                      |                        |                                      |                |
| Dose calculation                          | Х                       | Х                    | Х                      |                        |                                      |                |
| Review of final treatment plan            | Х                       | Х                    | Х                      | Х                      |                                      |                |

**Fig. 2** Roles and responsibilities of the radiation oncology team.<sup>6</sup>

to create and save templates of structure sets within the treatment-planning system (TPS) aids standardization and can be used to normalize practice between providers, within a health care system, or between institutions. In addition, some TPSs have a built-in anatomic atlas where contours of normal structures are displayed on a scrollable computed tomography (CT) scan. Sagittal/coronal planes and a 3-dimensional view can be used to review and adjust contours. Knowledge and application of recent advances can affect the efficiency and quality of OAR delineation. Artificial intelligence and deep learning are currently being applied to develop more sophisticated predictive models and automatic contouring software with the aim of improving the quality of treatment and the consistency of contours. 21 Auto-contouring features in the TPS can assist with multiple structures including the brain, lungs, spinal cord, and bony structures and may be reliable for certain structures (e.g., high-contrast organs such as femoral heads).

Reports have been published on the efficacy of autocontouring technology and the differences between automated and manual contouring.<sup>22,23</sup> Aselmaa et al focused on the quality of delineation and the efficiency within the workflow. The mean duration times required to contour a slice were captured for 3 cases. According to their review, quality was more consistent between physicians when automatic contouring technology was used. The study also confirmed the expectation in most cases regarding the time-saving aspect of this technology (Fig. 3).

Skitka et al evaluated the effects of automation bias on treatment planning, which pointed to possible errors associated with automation. This review noted that "participants with an aid made errors of omission (missed events when not explicitly prompted about them by the aid) and commission (did what an automated aid

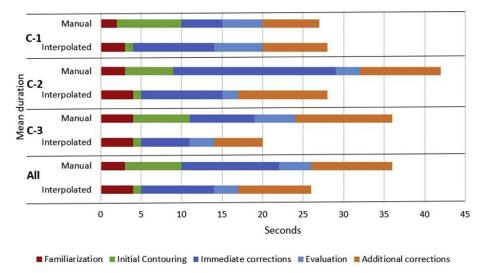
recommended, even when it contradicted their training and other 100% valid and available indicators)."<sup>23</sup>

These technology-reliant contouring tools promise to reduce time, improve consistency, and increase quality; however, as artificial intelligence is still under development in treatment planning, and contours generated from computer-based tools should always be subject to a careful manual review and adjustment before initiating treatment planning, and then formally reviewed as part of the plan approval process by the radiation oncologist.

# **Treatment Planning Processes**

#### Registration

Changes in anatomy due to possible tumor extension mandate a basic understanding of normal anatomy. To assist with delineating normal tissues, imaging considerations are also important. Certain OARs are better visualized or defined on contrast-enhanced CT scans or magnetic resonance imaging (MRI) scans, compared to CT scans alone. For example, it is easier to delineate the parotid gland on a contrast-enhanced CT due to the enhancing nature of the organ. Similarly, to accurately delineate the spinal cord, particularly for stereotactic planning, a high-resolution T2-weighted MRI or CT myelogram should be used. In the brain, it may be easier to delineate the optic chiasm and nerves using a fused high-resolution T1- or T2-weighted MRI rather than a typical treatment-planning CT. Contrast-enhanced CT scans, and specific sequences in MRI scans for contouring may facilitate delineation of normal tissues from adjacent tumor volumes and more adequately spare OARs compared with non-contrast treatment planning CT scans.



**Fig. 3** Mean durations (in seconds) of different workflow steps by slice in case C-1 (easy), C-2 (moderate difficulty), and C-3 (difficult). The type of workflow is labeled as manual or interpolated, meaning semi-automatic contouring. <sup>22</sup>
Reproduced with permission via the Creative commons license.

Image registration combines the information obtained from different imaging modalities with the planning CT and should be performed by a suitably trained member of the radiation oncology team.<sup>24</sup> During this process, special consideration must be taken as the patient position and morphology may differ between the diagnostic image and planning CT, resulting in potential uncertainties. As a result, alignment of images may require regional prioritization when variations do occur. All image registration, rigid or deformable, should be assessed for accuracy before contouring commences, including when autoregistration tools are used. Additionally, contours delineated using a registered image must be reviewed and verified on the planning CT before treatment planning commences and be evaluated during treatment plan approval by the radiation oncologist.

#### Segmentation

Contouring appropriate OARs is essential in the modern era of intensity modulated treatment planning, where specific details regarding tissue parameters are a necessary component of optimization and evaluation. Each organ has its own specific best practice(s) for appropriate contouring. For example, for lung constraints in the setting of definitive radiation therapy for lung cancer, tumor target volumes (eg, gross or clinical target volume) are usually subtracted from the OAR to assess volumetric dose constraints (eg, lung GTV).

Conversely, in the setting of central nervous system tumors, there can be no subtraction of target volumes because the entire organ must be assessed during the dose calculation. One approach when a target volume will be prescribed to a higher dose than the tolerance of an overlapping critical OAR is to create a structure that subtracts the critical OAR from the target (eg, PTV\_Eval). This may result in a compromise between adequate tissue sparing and optimal treatment of the target to maximize the therapeutic ratio of treatment.

In the development of guidance for appropriate organ contouring, the roles of artificial devices and surgical interventions in treatment planning were also reviewed. Recommendations to contour cardiac implantable electronic device<sup>25</sup> leads, ostomy bags,<sup>26</sup> and surgical hardware have been published. The presence of an implanted device may require additional consideration in treatment planning and a potential compromise between avoidance structures and optimal treatment plan design. This guidance does not specifically include these devices; however, they should be delineated when clinically appropriate.

In other settings, a planning OAR volume (PRV) provides a margin around an OAR to account for variations in setup, motion, and anatomy when appropriate.<sup>27</sup> A PRV will further ensure the sparing of particularly sensitive OARs, or better shape dose distributions by intentionally

constraining an area to a lower dose. This use of a PRV is most common in the treatment of tumors close to critical neural structures (eg, spinal cord or optic structures) to prevent unintentional overdosing of these OARs owing to setup errors (particularly with complex treatment planning techniques). In less complex cases, the spinal canal is commonly used as a surrogate for the spinal cord.

# Treatment plan optimization and evaluation

Incorrect contours may result in difficulties with plan optimization for inverse planning or multileaf collimator placement in forward-planned techniques. Ultimately, an error in any planning phase may have a downstream consequence that results in a suboptimal plan producing suboptimal patient outcomes.

In accepting radiation therapy plans, a key step in the process is plan review and 3-dimensional dose-volume outcome analysis. Whether a more meaningful outcome for an OAR is the mean, maximum, or volumetric dose, inherent in the interpretation of that metric is the assumption that the contours represent the OAR. If not, the entire dosimetric treatment plan may be compromised because treatment design aspects including beam angle, location of static beams, and avoidance regions in arcs may be affected. By using a standard set of tissues contoured based on disease site, DVH data can be analyzed confidently.

Some universal suggestions are relevant regardless of disease site. This list is not meant to be exhaustive, but represents some general guiding principles:

- Contour on appropriate density windows (i.e., bone, lung) for each tissue.
- Create structure set templates in the TPS to set standard practice procedures.
- Use contrast-enhanced scans when relevant to assist
  with tissue delineation. Large areas of contrast may
  need a density override or a registered noncontrast
  scan for planning purposes if volumes of contrast
  significantly influence dose calculations.
- Register relevant imaging (eg, MRI, contrastenhanced CT) to the planning CT to assist with tissue definition, but always review the final contouring on the primary data set. Ascertain the best region of alignment when there are inconsistencies between the images.
- Review structures in the coronal and sagittal planes when contouring on axial slices to verify completeness of coverage in all dimensions.

# Summary

Standardization improves patient safety, efficiency, and accuracy in radiation oncology treatment. This consensus guidance represents an ASTRO-quality

initiative to provide recommendations for the standardization of normal-tissue contouring that is performed during external beam treatment planning for each anatomic treatment site. Table E1 defines 2 sets of structures for anatomic sites: those that are recommended in all adult definitive cases and may assist with organ selection for palliative cases, and those that should be considered on a case-by-case basis depending on the specific clinical scenario. Table E2 outlines some resources available to define the parameters of general OAR tissue delineation. Using this paper in conjunction with resources that define tissue delineation parameters and published dose constraints will enable practices to develop a consistent approach to normal-tissue evaluation and dose documentation.

# **Acknowledgments**

This document is endorsed by American Association of Medical Dosimetrists. The authors gratefully acknowledge the peer reviewers: Jennifer Bellon, MD, Nicolas DeNunzio, MD, MS, Avraham Eisbruch, MD, Supriya Jain, MD, Lori Kasuski, RT(R)(T), CMD, Aaron Kusano, MD, Lilie Lin, MD, Simon Lo, MD, Billy Loo, MD, Charles Mayo, PhD, Sasa Mutic, PhD, Tracey Schefter, MD, Abhishek Solanski, MD, Stephanie Weiss, MD, MS, Ying Xiao, PhD, Catheryn Yashar, MD, Sandra Zaky, MD, and Michael Zelefsky, MD. This document was prepared by the American Society for Radiation Oncology (ASTRO) as part of an ongoing quality initiative with the goal of enabling members to consistently deliver the highest quality and value care to cancer patients.

# **Supplementary Data**

Supplementary material for this article can be found at https://doi.org/10.1016/j.prro.2018.12.003.<sup>28-48</sup>

#### References

- Emami B, Lyman J, Brown A, et al. Tolerance of normal tissue to therapeutic irradiation. *Int J Radiat Oncol Biol Phys.* 1991;21:109-122.
- Benedict SH, Yenice KM, Followill D, et al. Stereotactic body radiation therapy: the report of AAPM Task Group 101. *Med Phys*. 2010;37:4078-4101.
- Bentzen SM, Constine LS, Deasy JO, et al. Quantitative Analyses of Normal Tissue Effects in the Clinic (QUANTEC): an introduction to the scientific issues. *Int J Radiat Oncol Biol Phys.* 2010;76(3 Suppl): \$3.50
- Moran JM, Molineu A, Kruse JJ, et al. Executive summary of AAPM Report Task Group 113: Guidance for the physics aspects of clinical trials. J Appl Clin Med Phys. 2018.
- Mayo CS, Moran JM, Bosch W, et al. American Association of Physicists in Medicine Task Group 263: Standardizing

- Nomenclatures in Radiation Oncology. *Int J Radiat Oncol Biol Phys.* 2018;100:1057-1066.
- Zietman A, Palta JR, Steinburg ML, Blumberg AL, Burns RA, Cagle SW. Safety is no accident: A framework for quality radiation oncology and care. Fairfax, VA: ASTRO; 2012.
- Cattaneo GM, Reni M, Rizzo G, et al. Target delineation in postoperative radiotherapy of brain gliomas: interobserver variability and impact of image registration of MR(pre-operative) images on treatment planning CT scans. *Radiother Oncol.* 2005;75:217-223.
- Yamamoto M, Nagata Y, Okajima K, et al. Differences in target outline delineation from CT scans of brain tumours using different methods and different observers. *Radiother Oncol*. 1999;50:151-156
- Hall WH, Guiou M, Lee NY, et al. Development and validation of a standardized method for contouring the brachial plexus: preliminary dosimetric analysis among patients treated with IMRT for head-andneck cancer. *Int J Radiat Oncol Biol Phys.* 2008;72:1362-1367.
- Nelms BE, Robinson G, Markham J, et al. Variation in external beam treatment plan quality: An inter-institutional study of planners and planning systems. *Pract Radiat Oncol.* 2012;2:296-305.
- Spoelstra FO, Senan S, Le Péchoux C, et al. Variations in target volume definition for postoperative radiotherapy in stage III nonsmall-cell lung cancer: analysis of an international contouring study. *Int J Radiat Oncol Biol Phys.* 2010;76:1106-1113.
- Li XA, Tai A, Arthur DW, 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;73:944-951.
- Van de Steene J, Linthout N, de Mey J, et al. Definition of gross tumor volume in lung cancer: inter-observer variability. *Radiother Oncol*. 2002;62:37-49.
- Rosewall T, Bayley AJ, Chung P, et al. The effect of delineation method and observer variability on bladder dose-volume histograms for prostate intensity modulated radiotherapy. *Radiother Oncol*. 2011:101:479-485.
- 15. Foppiano F, Fiorino C, Frezza G, Greco C, Valdagni R. AIRO National Working Group on Prostate Radiotherapy. The impact of contouring uncertainty on rectal 3D dose-volume data: results of a dummy run in a multicenter trial (AIROPROS01-02). *Int J Radiat Oncol Biol Phys.* 2003;57:573-579.
- Nelms BE, Tomé WA, Robinson G, Wheeler J. Variations in the contouring of organs at risk: test case from a patient with oropharyngeal cancer. *Int J Radiat Oncol Biol Phys.* 2012;82:368-378.
- PSO, C. Aggregate Report Patients Safety Work Product Q4 2017. 2018.
- Group, P.S.I.R.S. Radiotherapy Errors and Near Misses Data Report (December 2013 to November 2015). Public Health England; 2016.
- ASTRO. APEx Program Standards; 2016 [cited 2016]; Available from https://www.astro.org/uploadedFiles/\_MAIN\_SITE/Daily\_Prac tice/Accreditation/Content\_Pieces/ProgramStandards.pdf.
- Scope of Practice of a Medical Dosimetrist. American Association of Medical Dosimetrists; 2012.
- Kearney V. The application of artificial intelligence in the IMRT planning process for head and neck cancer. *Oral Oncol.* 2018;87: 111-116.
- Aselmaa A, van Herk M, Song Y, Goossens RHM, Laprie A. The influence of automation on tumor contouring. *Cognition, Technol*ogy & Work. 2017;19:795-808.
- Skitka LJ, Mosier KL, Burdick M. Does automation bias decisionmaking? *International Journal of Human Computer Studies*. 1999; 51:991-1006.
- Brock KK, Mutic S, McNutt TR, Li H, Kessler M. Use of image registration and fusion algorithms and techniques in radiotherapy: Report of the AAPM Radiation Therapy Committee Task Group No. 132. *Med Phys.* 2017;44:e43-e76.
- Indik JH, Gimbel JR, Abe H, et al. 2017 HRS expert consensus statement on magnetic resonance imaging and radiation exposure in

- patients with cardiovascular implantable electronic devices. *Heart Rhythm.* 2017;14:e97-e153.
- 26. Baumann BC, Bosch WR, Bahl A, et al. Development and Validation of Consensus Contouring Guidelines for Adjuvant Radiation Therapy for Bladder Cancer After Radical Cystectomy. *Int J Radiat Oncol Biol Phys.* 2016;96:78-86.
- Hodapp N. [The ICRU Report 83: prescribing, recording and reporting photon-beam intensity-modulated radiation therapy (IMRT)]. Strahlenther Onkol. 2012;188:97-99.
- 28. Brouwer CL, Steenbakkers RJ, Bourhis J, et al. CT-based delineation of organs at risk in the head and neck region: DAHANCA, EORTC, GORTEC, HKNPCSG, NCIC CTG, NCRI, NRG Oncology and TROG consensus guidelines. *Radiother Oncol.* 2015; 117:83-90.
- Balagamwala EH. Stereotactic body radiotherapy for the treatment of spinal metasteses. *Journal of Radiation Oncology*. 2012;1:255-265.
- Gondi V, Tome WA, Rowley H.A, Mehta M.P., Hippocampal Contouring: A Contouring Atlas for RTOG 9333.
- Kao J, Darakchiev B, Conboy L, et al. Tumor Directed, Scalp Sparing Intensity Modulated Whole Brain Radiotherapy for Brain Metastases. *Technol Cancer Res Treat*. 2015;14:547-555.
- Scoccianti S, Detti B, Gadda D, et al. Organs at risk in the brain and their dose-constraints in adults and in children: a radiation oncologist's guide for delineation in everyday practice. *Radiother Oncol*. 2015;114: 230-238.
- Pacholke HD, Amdur RJ, Schmalfuss IM, Louis D, Mendenhall WM. Contouring the middle and inner ear on radiotherapy planning scans. Am J Clin Oncol. 2005;28:143-147.
- 34. Feng FY, Kim HM, Lyden TH, et al. Intensity-modulated radiotherapy of head and neck cancer aiming to reduce dysphagia: early dose-effect relationships for the swallowing structures. *Int J Radiat Oncol Biol Phys.* 2007;68:1289-1298.
- Christianen ME, Langendijk JA, Westerlaan HE, van de Water TA, Bijl HP. Delineation of organs at risk involved in swallowing for radiotherapy treatment planning. *Radiother Oncol*. 2011;101:394-402
- Rosenthal DI, Fuller CD, Barker JL Jr, et al. Simple carotid-sparing intensity-modulated radiotherapy technique and preliminary experience for T1-2 glottic cancer. *Int J Radiat Oncol Biol Phys.* 2010; 77:455-461.
- Freedman L, Sidani C. A radiation oncologist's guide to contour the parotid gland. *Pract Radiat Oncol*. 2016;6:e315-e317.

- 38. Kong FM, Ritter T, Quint DJ, et al. Consideration of dose limits for organs at risk of thoracic radiotherapy: atlas for lung, proximal bronchial tree, esophagus, spinal cord, ribs, and brachial plexus. *Int J Radiat Oncol Biol Phys.* 2011;81:1442-1457.
- 39. Nielsen MH, Berg M, Pedersen AN, et al. Delineation of target volumes and organs at risk in adjuvant radiotherapy of early breast cancer: national guidelines and contouring atlas by the Danish Breast Cancer Cooperative Group. Acta Oncol. 2013;52:703-710.
- 40. White J, T.A., Arthur D, Buchholz T, MacDonald S, Marks L, Pierce L, Recht A, Rabinovitch R, Taghian A, Vicini F, Woodward W, Li X.A., NRG Breast Cancer Atlas for Radiation Therapy Planning: Consensus Definition.
- Feng M, Moran JM, Koelling T, et al. Development and validation of a heart atlas to study cardiac exposure to radiation following treatment for breast cancer. *Int J Radiat Oncol Biol Phys.* 2011;79: 10-18
- Duane F, Aznar MC, Bartlett F, et al. A cardiac contouring atlas for radiotherapy. *Radiother Oncol*. 2017;122:416-422.
- 43. Lorenzen EL, Taylor CW, Maraldo M, et al. Inter-observer variation in delineation of the heart and left anterior descending coronary artery in radiotherapy for breast cancer: a multi-centre study from Denmark and the UK. *Radiother Oncol.* 2013;108:254-258.
- 44. Lee J, Hua KL, Hsu SM, et al. Development of delineation for the left anterior descending coronary artery region in left breast cancer radiotherapy: An optimized organ at risk. *Radiother Oncol*. 2017;122:423-430.
- Jabbour SK, Hashem SA, Bosch W, et al. Upper abdominal normal organ contouring guidelines and atlas: a Radiation Therapy Oncology Group consensus. *Pract Radiat Oncol.* 2014;4:82-89.
- Brooks C, Hansen VN, Riddell A, Harris VA, Tait DM. Proposed genitalia contouring guidelines in anal cancer intensity-modulated radiotherapy. *Br J Radiol*. 2015;88:20150032.
- Gay HA, Barthold HJ, O'Meara E, et al. Pelvic normal tissue contouring guidelines for radiation therapy: a Radiation Therapy Oncology Group consensus panel atlas. *Int J Radiat Oncol Biol Phys.* 2012;83:e353-e362.
- 48. Yi SK, Mak W, Yang CC, et al. Development of a standardized method for contouring the lumbosacral plexus: a preliminary dosimetric analysis of this organ at risk among 15 patients treated with intensity-modulated radiotherapy for lower gastrointestinal cancers and the incidence of radiation-induced lumbosacral plexopathy. *Int J Radiat Oncol Biol Phys.* 2012;84:376-382.