Global Advanced Research Journal of Medicine and Medical Science (ISSN: 2315-5159) Vol. 3(5) pp. 090-094, May 2014 Available online http://garj.org/garjmms/index.htm Copyright © 2014 Global Advanced Research Journals

Full Length Research Paper

Calculation of organs radiation dose in cervical carcinoma external irradiation beam using day's methods

Yousif M. Yousif Abdallah^{1*,} Mohamed E. Gar-elnabi¹, Abdoelrahman H. A. Bakary¹, Alaa M. H. Eltoum¹, Abdelazeem K. M. Ali²

¹Radiotherapy and Nuclear Medicine Department, College of Medical Radiological Science, Sudan University of Science and Technology, Khartoum, Sudan
²Radiation Oncology Department, National Cancer Institute, University of Gazeria, Madani, Sudan

Accepted 19 March, 2014

The study was established to measure the amount of radiation outside the treatment field in external beam radiation therapy using day method of dose calculation, the data was collected from 89 patients of cervical carcinoma in order to determine if the dose outside side the irradiation treatment field for spleen, liver, both kidneys, small bowel, large colon, skin within the acceptable limit or not. The cervical field included mainly 4 organs which are bladder, rectum part of small bowel and hip joint these organ received mean dose of (4781.987±281.321), (4736.91±331.8), (4647.64±387.1) and (4745.91±321.11) respectively. The mean dose received by outfield organs was (77.69±15.24cGy) to large colon, (93.079±12.31cGy) to right kidney (80.688±12.644cGy) to skin, (155.86±17.69cGy) to small bowel. This was more significant value noted.

Keywords: Radiation Dose, Cervical Carcinoma, Day's Methods

INTRODUCTION

In 2008 it was estimated that 529 000 incident cases and 275 000 deaths due to carcinoma of the uterine cervix (cervical cancer) occurred annually worldwide. About 88% of this burden is borne by low and middle income countries (LMC) where cervical cancer is the leading malignancy among women (Day, 1950; Eifel et al., 2004). Screening with Pap smear decreases mortality by 70%. The mean age of women diagnosed with cervical intraepithelial neoplasia (CIN) is 15–20 years younger than those diagnosed with invasive disease. ACS

recommends screening for all women who are sexually active or >20 years old. Following three normal annual exams after age 30, screening may be performed less frequently, at least once every 3 years (From clinical trials to clinical practice). The associated risk factors: early first intercourse, multiple partners, history of other STD's, high parity, smoking, immunosuppression, and prenatal DES exposure (clear cell CA). With 90-95% of cases is associated with HPV infection. More types 16 and 18 confer the highest risk of SCC and adenocarcinoma, respectively. HPV 6 and 11 are associated with benign warts. 80-90% of invasive tumors are SCC, 10–20% is adenocarcinoma, and 1–2% is clear cell. Preinvasive disease include the ASCUS (2/3 resolve spontaneously. Repeat Pap in 6 months and, if abnormal, perform

^{*}Corresponding Author E-mail: yousifmohamed@sustech.edu, yousifmohamed@outlook.com

colposcopy), LGSIL, and HGSIL. Prognostic factors include LN metastases, tumor size, stage, uterine extension, and Hgb level <10. With the risk of pelvic LN involvement for stage I, II, and III disease is approximately 15%, 30%, and 45%, respectively (Haie-Meder et al., 2005). Such cancerous disease can be diagnosed by Pap smear if not bleeding. Colposcopy, Cystoscopy, sigmoidoscopy, and/or barium enema for IIB, III, or IVA disease, or for symptoms, Laboratory tests and Imaging with CT/MRI of abdomen and pelvis and CXR. PET scans are sensitive (~85–90%) and specific (~95–100%). If stage IIIB, place renal stent prior to starting chemotherapy (Hansen et al., 2010).

In this realm several calculations is carried out carefully to determine level of doses out the field limits. EBPT is unavoidably associated with irradiation, at lower doses, of large volumes of normal tissue away from the beam path (Jane et al., 2009; Jeffrey et al., 2010; Johnsson et al., 1997; Keys et al., 1997). According to the latest recommendations of (ICRU) concerning the remaining volume at risk (RVR), the search for means of more accurately determining such doses is of renewed clinical interest. Indeed, according to ICRU Report 83 (ICRU 2010), all normal tissues that could potentially be irradiated should be included in the RVR, and the absorbed dose in the RVR might be useful for estimating risk of later effects such as carcinogenesis. In essence, the out-of-field dose arises from three main sources: (1) leakage from the treatment unit; (2) scatter from the treatment unit head and from beam modifiers such as wedges and blocks; and (3) internal scatter originating in the patient. Different scientists estimated the dose to points in the body outside the primary beam. Therefore a generalized model is developed to calculate this dose with reasonable accuracy better than ±30% (Keys et al., 1999; Keys et al., 2003; Kim et al., 2008). Radiation scattered in the patient and the radiation scattered from the collimator exhibit a strong dependence on field size and distance and are predominant only at short distances. At larger distances large amount of leakage with accuracy is better than ±50. Measurement of peripheral dose (PD), for instance, to the gonads, for treatment machines and/or techniques. specific Published data were available for ⁶⁰CO, 4, 6, 8, and 10 MV, and 18 to 25 for a large verity of treatment machines. Furthermore, an analysis of possible corrections for depth dependence, field elongation, irregularly shaped fields, wedges, and shielding blocks which affect received dose (Landoni et al., 1997). Some occasion when it measurement of dose level outside of field is proves to give radiotherapy to a pregnant patient. Especially at the time when pregnancy has not been confirmed, levels of radiation dose. The Code of Practice for the Protection of Persons against Ionizing Radiations provides that an occupationally exposed female should not receive in excess of 1.3rem, i.e. 0.013Gy, to the abdomen. Thus a maximum occupational exposure of 0.023Gy is "accepted" by the code of practice.

The characterization of the incident photon beam is usually divided into its dependence on collimator setting (head-scatter factor) and off-axis position (primary offaxis ratio) (Howell et al., 2010). These parameters are normally measured "in air" with a build-up cap thick enough to generate full dose build-up at the depth of dose maximum. Unwanted radiation has been measured as a function of the distance outside the primary beam, and field size because this absorbed dose outside the radiation fields is clinically important, potentially affecting cataract formation, gonadal function, and fertility (Rafi et al., 2003). This dose can also be responsible for exposure to the fetus in a pregnant woman, and dose to breast and carcinogenesis may be a concern. Using a locally fabricated water phantom of dimensions 45cm×45cm×30 cm at 5.0 cm depth for horizontal beam. In the present study, a 0.1 cc ion chamber type 23323 in conjunction with a PTW UNIDOS electrometer has been used for dose measurement. Collimator-related radiation dose was about 3 times higher than that from the more modern machine. Therefore the scattered and leakage radiation show a strong dependence on field size and distance to the beam axis and is predominant only at short distances (Mohamed et al., 2012; Morris et al., 1999; Pearcey et al., 2002). In-field radiation doses can be accurately and rapidly calculated using commercially available treatment planning systems (TPSs) (Peters et al., 2000: Rose et al., 1999). These TPSs do not. however, accurately model doses outside the treatment field, nor are they commissioned for such calculations. A recent study evaluated the accuracy with which a commercial TPS calculated absorbed dose in regions where the isodose lines reported by the TPS were less than 5% of the prescribed dose (Rotman et al., 1995; Rotman et al., 2006). Which demonstrated that in this very low stray dose region, the predicted doses were at worst 60% lower than corresponding measured data and that the accuracy of the TPS calculated doses decreased with increasing distance from the treatment field. In CPRT, out-of-field organs are easily defined by their proximity to the field border which is defined by the collimating jaws. Radiation dose measurements in anthropomorphic phantoms are considered the gold standard in peripheral dose assessment and have frequently been used to determine peripheral organ doses in studies of radiation-induced late effects from photon radiotherapy (Stehman et al., 2007). In range of 3.75-11.25 cm from the edge of the treatment field, the TPS underestimated dose by an average of 40% ± 20%. As the distance from the treatment field increased, the TPS underestimated the dose with increasing magnitude. radiation Documents dosage sensitive to organs/structures located outside the radiotherapeutic target volume for four treatment situations: (a) head and

neck, (b) brain (pituitary and temporal lobe), (c) breast and (d) pelvis.

MATERIALS AND METHODS

The data of this study was collected from NCI Aljazeera state with patients treated from cervical cancer radically using external radiation therapy. The treatment was delivered using two opposed fields 'anterior and posterior field' encompasses the true pelvis including the cervix, bladder and rectum. As well as uterus, parametrial tissue, part of small bowel and hip joint. The data were collected from the patient included the field sizes, AP separation, lateral separation, tumor dose, patient weight, height, given dose, Then from the patient CT images the distance from the field border with antro-posterior and postro-anterior depths of transfers colon, both kidneys, liver, skin depth at certain distance from the field border in addition to small bowel at outfield region, spleen depths and cervix as well were measured. The dose received by these organ from each field was calculated collectively using central dose calculation in respect to its positions from the mid-depth 'separation' of the Anterior and posterior field using day's method outside the field in respect to their arbitrary center.

Day's method for dose calculation outside the irradiated field

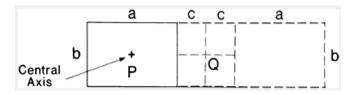


Figure 1. Shows Calculation of depth dose outside a rectangular field

The distances of the critical organs, then the % dose received by the critical were calculated using day's method as follows; suppose Q is a point outside the field at a distance c from the field border. Imagine a rectangle adjacent to the field such that it contains point Q and has dimensions 2c. Place another rectangle of dimensions a b on the other side of Q such that the field on the right of Q is a mirror image of the field on the left, as shown in the figure. The dose at point Q at depth d is then given by subtracting the depth dose at Q for field $2c \times b$ from that for field $(2a + 2c) \times b$ and dividing by 2. The procedure is

illustrated by the following example. Suppose it is required to determine percent depth dose at Q (relative to D_{max} at P) outside a 15 ×10 cm field at a distance of 5 cm from the field border. In Fig. 1 then, a = 15, b = 10, and c = 5. Suppose Q is at the center of the middle rectangle of dimensions $2c \times b$. Then the dose D_Q at 10-cm depth is given by: $\frac{1}{2}[D_Q(40 \times 10) - D_Q(10 \times 10)]$

If D_Q is normalized to D_{max} at P, one gets the percent depth dose at Q or $\%D_Q$.

$$\%D_Q = \frac{1}{BSF(15 \times 15)} \times \frac{1}{2} [BSF(40 \times 10) \times \%DD(40 \times 10) - 1]$$

Thus for a ⁶⁰Co beam at $SSD = 80$ cm.

$$\%D_Q = \frac{1}{(1.052)} \times \frac{1}{2} [1.054 \times 58.8 - 1.036 \times 55.6] = 2.1$$

RESULTS

Table 1. show (mean ±Std. deviation) of dose (cGy) received by organs outside the radiation treatment field in treatment 50 patient of cervical cancer

Organs	Min (cGy)	Max (cGy)	Mean ±Std. Deviation
Large bowel	48.1	128.5	77.7±15.2
Left kidney	53.8	113.4	74.8±11.6
Liver	44.8	95.7	62±9.6
Right kidney	68.7	137.2	93 ±12.3
Skin	51.9	119.7	80.7±12.6
Small bowel	117.5	210.6	155.9±17.7
Spleen	42.1	87.3	56.9±7.5

Table 2. Shows Mean ±Std. Deviation of the parameters used in dose calculation for cervical cancer Patient weight is 61.3±12.2, height 163.3±6.8, given dose 8409.5±600.1 and patient separation was 18.9±1.6

Organ	AP depth (cm)	PA depth (cm)	Distance from field border(cm)
Small bowel	5.7±0.35	13.22±1.244	6.384±0.117
Large colon	2.95±1.033	16.84±0.88	13.6±1.57
Liver	10.64±.398	8.28±1.19	16.96±1.31
Spleen	11.3±1.42	7.584±0.17	18.07±0.67
Skin	18.176±1.2	14.8±0.16	2.00±.000
Left kidney	11.16±1.28	7.79±0.321	14.44±1.25
Right kidney	11.8±1.3	7.2±0.28	11.6±0.9

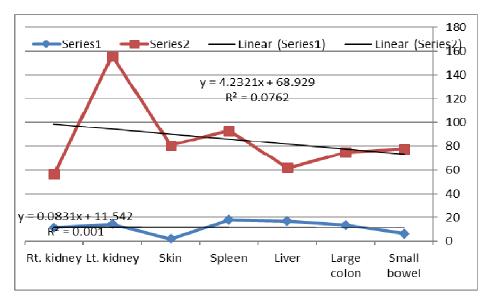


Figure 2. An illustration of radiation map created for doses received by organ inside and outside field margin of cervical cancer as distance from it measured by cm (arrowed).

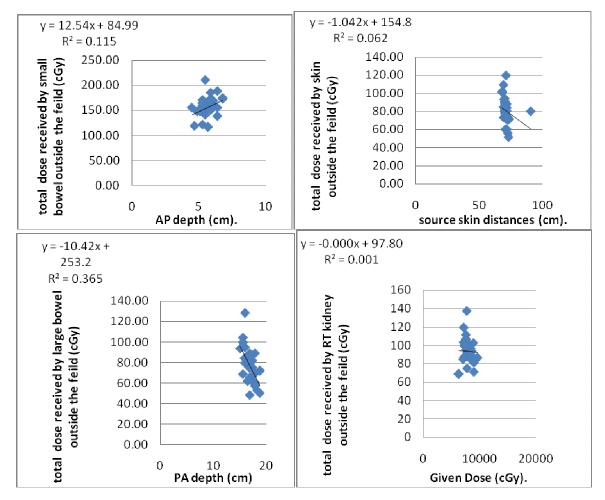


Figure 3. A, B, C, D, E Shows total dose received by small bowel, skin, large bowel and Right kidney.

CONCLUSION

The study were established to measurement the amount of radiation outside the treatment field in external beam radiation therapy using day method of dose calculation, the data were collected from 89 patients of cervical carcinoma in order to determine if the dose outside side the irradiation treatment field for spleen, liver, both kidneys, small bowel, large colon, skin within the acceptable limit or not. The method for assessing organ doses throughout the body from photon radiotherapy described here can be used in studies that require accurate knowledge of a wide range of doses from both primary and scatter radiation, especially the scatter radiation which it is contribution consider to be very critical issue in EBRT, so the quality assurance test should be carried out to assess the amount of leakage radiation and scatter radiation outside of definite field size to determine if desirable radiotherapy dose distribution and calculation design for specific treatment field can delivered the radiation with high therapeutic ratio. Such broad information will be of particular use in studies of radiation-induced late effects, which require accurate knowledge of doses to in-field, out of-field and partially infield organs to predict the risk to organs throughout the body.

The mean dose received by outfield organs was (77.69 \pm 15.24cGy) to large colon, (93.079 \pm 12.31cGy) to right kidney (80.688 \pm 12.644cGy) to skin, (155.86 \pm 17.69cGy) to small bowel. This was more significant value noted.

REFERENCES

- Almberg SS (2012). Monte Carlo study of in-field and out-of-field dose distributions from a linear accelerator operating with and without a flattening-filter. J. med. phys. 39 (10).
- Day MI (1950). A note on the calculation of dose in x-ray fields. Br. J. Radiol. 23:368.
- Eifel PJ, Winter K, Morris M, et al (2004). Pelvic irradiation with concurrent chemotherapy versus pelvic and para-aortic irradiation for high-risk cervical cancer: an update of radiation therapy oncology group trial (ROG) 90-01. J. Clin. Oncol. 22:872-880.
- From clinical trials to clinical practice; a user's guide to evidence-based oncology, Eur. J. Cancer 6 93.
- Haie-Meder C, Pötter R, Van Limbergen E, et al (2005). Recommendations from Gynaecological (GYN) GEC-ESTRO Working Group (I): concepts and terms in 3D image based 3D treatment planning in cervix cancer brachytherapy with emphasis on MRI assessment of GTV and CTV. Radiother .Oncol.74(3):235-245.
- Hansen K et al (2010). Handbook of Evidence-Based Radiation Oncology ,2nd Edition, Springer, New York, p 499-511.
- Howell MR, et al (2010). Methodology for determining doses to in-field, out-of-field and partially in-field organs for late effects studies in photon radiotherapy. J. physics in med. biol. 55: 7009–7023.
- Jane De et al (2009). Practical Radiotherapy Planning, 4rd Edition, Arnod Publisher, Lodon p.
- Jeffrey A, et al (2010). Clinical Radiation Oncology, third edition, Elsevier Inc., Philadelphia, p 1037-1177
- Johnsson SA, et al (1997). Off-axis primary-dose measurements using a mini-phantom. J. Med. Physics. 24: 763-767.

- Keys HM, Bundy BN, Stehman FB, et al (1997). Adjuvant hysterectomy after radiation therapy reduces detection of local recurrences in "bulky" stage IB cervical without improving survival: results of a prospective randomized GOG trial. Cancer J. Sci. Am. 3:117(abstr).
- Keys HM, Bundy BN, Stehman FB, et al (1999). Cisplatin, radiation, and adjuvant hysterectomy compared with radiation and adjuvant hysterectomy for bulky stage IB cervical carcinoma. N. Engl. J. Med. 340:1154-1161.
- Keys HM, Bundy BN, Stehman FB, et al (2003). Radiation therapy with and without extrafascial hysterectomy for bulky stage IB cervical carcinoma: a randomized trial of the Gynecologic Oncology Group. Gynecol. Oncol. 89:343-353.
- Kim HJ, et al (2008). Are doses to ICRU reference points valuable for predicting late rectal and bladder morbidity after definitive radiotherapy in uterine cervix cancer? J. Tumori. 94: 327-332
- Landoni F, Maneo A, Colombo A, et al (1997). Randomised study of radical surgery versus radiotherapy for stage lb-lla cervical cancer. Lancet. 350:535-540.
- Mohamed A, et al (2012). A multi-plane source model for out-of-field head scatter dose calculations in external beam photon therapy. J. physics in med. biol. 57.
- Morris M, Eifel PJ, Lu J, et al (1999). Pelvic radiation with concurrent chemotherapy compared with pelvic and para-aortic radiation for high-risk cervical cancer. N. Engl. J. Med. 340: 1137-1143.
- Pearcey R, Brundage M, Drouin P, et al (2002). Phase III trial comparing radical radiotherapy with and without cisplatin chemotherapy in patients with advanced squamous cell cancer of the cervix. J. Clin. Oncol. 20:966-972.
- Peters WA III, Liu PY, Barrett RJ II, et al (2000). Concurrent chemotherapy and pelvic radiation therapy compared with pelvic radiation therapy alone as adjuvant therapy after radical surgery in high-risk early-stage cancer of the cervix. J. Clin. Oncol. 18:1606-1613.
- Rafi M Uddin, et al (2003) MEASUREMENT OF DOSE OUT SIDE THE IRRADIATED VOLUME BY USING LOCALLY FABRICATED WATER PHANTOM, Bangladesh J. Med. Physics 2: 41-43
- Rose PG, Bundy BN, Watkins EB, et al (1999). Concurrent cisplatin-based radiotherapy and chemotherapy for locally advanced cervical cancer. N. Engl. J. Med. 340:1144-1153.
- Rotman M, Pajak TF, Choi K, et al (1995). Prophylactic extended-field irradiation of para-aortic lymph nodes in stages IIB and bulky IB and IIA cervical carcinomas. Ten-year treatment results of RTOG 79-20. JAMA. 274:387-393.
- Rotman M, Sedlis A, Piedmonte MR, et al (2006). A phase III randomized trial of postoperative pelvic irradiation in stage IB cervical carcinoma with poor prognostic features: follow-up of agynecologic oncology group study. Int. J. Radiat. Oncol. Biol. Phys. 65:169-176.
- SIR (1984). Dose levels outside radiotherapy beams. The Br. J. Radiol. 57: 274-275
- Stehman FB, Ali S, Keys HM, et al (2007). Radiation therapy with or without weekly cisplatin for bulky stage 1B cervical carcinoma: follow-up of a Gynecologic Oncology Group trial. Am. J. Obstet. Gynecol. 197(5):503.e1-6.
- Stovall M (2010). Tissue doses from radiotherapy of cancer of the uterine cervix. J. cancer res. theraputric. 6: 482- 486
- Van Der Giessen (1996). Collimator-related radiation dose for different cobalt machines and linear accelerators. Int. J. Radiation Oncol. *Biol. *Phys. 35: 399–405
- VanDer Giessen PH, et al (1994). Calculation and measurement of the dose at points outside the primary beam for photon energies of 6, 10, and 23 MV. Int. j. radiat. oncol. phys. biol. 30(5): 1239–1246
- VanDer Giessen, Coen WH (1993). Calculation and measurement of the dose to points outside the primary beam for CO-60 gamma radiation. Int. J. Radiat. Oncol. * Biol. * Phys. 27(3): 717–724
- VanDer.Giessen PH (1996). A simple and generally applicable method to estimate the peripheral dose in radiation teletherapy with high energy X-rays or gamma radiation. Int. j. rad. oncol. phys. biol. 35: 1059–1068