# Prediction of Dose Calculation of Breast and Chest Tumors Using Different Algorithms

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This work significantly observes the suitable of algorithms by comparing the dose calculation of breast and chest tumors using Clarkson, Convolution, Superposition and Fast Superposition algorithms in three-dimensional Conformal Radiotherapy (3D-CRT). Thirteen post-operative breast cases and fifteen cancer chest patients were taken in this work. Treatment plans were created using 6 and 15 MV Photon beam quality with the CMS XiO treatment planning system (Computerized Medical System, St. Louis, MO). Statistical analysis was done by relating the mean relative difference, conformity index, and homogeneity index for target structures. The percentage of difference noted between algorithms were 9.4%, recorded with a low density point in case of breast. Superposition and fast superposition algorithm were presented very good results in both cases in view of the mean relative variances using a prescribed dose with four algorithms and the lowest relative variable using a given dose in different treatment techniques PTV. Superposition algorithm exposed good calculations in all techniques. Clarkson algorithm showed monotonically variation in the dose calculation points and histogram parameters. The other three algorithms are shown approximately similar results. We found from the dosimetric results a considerable difference among four algorithms. Conferring to this study, the algorithms varied. Significant upkeep must be occupied when calculating treatment plans, as the selection of the dose computation algorithm may affected on treatment planning as well as clinical effects.

Keywords: Algorithm, chest tumors, cancer breast, homogeneity index, treatment planning system.

The radiotherapy is useful to provide a given dose to tumor specifically while minimizing dose to critical organs. The radiation dose is a basis of the process, which also includes dose computation and delivery of radiation beam. The previous is the main section of a treatment planning system. Accuracy and features of any treatment planning system (TPS) are dependent on the type of algorithms which used in steps of the planning procedure. An algorithm is defined as a sequence of instructions that operates on a set of input data, converting that information into a set of ultimate results that are of interest to the user. Many algorithms are used in the treatment planning

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progression. The most recognized dose calculation algorithm that generates the dose at any point in the patient while taking into account beam characteristics and the patient<sup>1</sup>. It is a principal importance for the present conformal radiotherapy technique to have accuracy in dose computations in almost all applicable clinical situations. Treatment of lung tumors is One of these situations where irradiation has to be planned under stimulating conditions for dose calculation<sup>2</sup>. The patient dose predictions accuracy has continuously improved by moving from the simple scatter inhomogeneity corrections over pencil beam algorithms to point kernel-based Convolution/ Superposition methods<sup>3</sup>. In our study Clarkson, convolution, superposition, and fast superposition algorithms were applied to all plans. Our aim of this study to compare the results from four algorithms, Six dissimilar points, demonstrating different heterogeneity conditions and position using 3D-CRT"Three-Dimensional Conformal Radiotherapy (3D-CRT)" Technique. This helpful for knowing the suitability of an algorithm for the particular diagnostic and treatment technique. Conformity Index, Homogeneity Index, Mean Dose, and Mean Relative Difference have been used to evaluate the external beam plans. Also, Dose Volume Histograms (DVH) for different structures were calculated to quantify the dose to the other OARs.

#### **MATERIALSAND METHODS**

## **Patients and techniques**

Twenty eight Cancer patients were examined for two treatments position in the chest region: thirteen breast cases tangential 6,15 MV photons to the chest wall with 2-4 fields for treatment, The other fifteen patient inside chest wall treated with 2-6 fields with 6,15 MV photon beam. Doses of 200 cGy/Fraction, were set to the planning target volume of the breast and other chest legion. Planning Target Volume (PTV) was getting drawn using the 5mm isotropic extension of the Clinical Target Volume (CTV), which in turn was gained from the macroscopic Gross Tumor Volume (GTV). The PTV was made to eliminate other OARs by 5mm<sup>4</sup>. Treatment planning designs for the target and OARs were created using 6MV, 15 MV and mixed 6 and 15 MV Photon beam quality using Clarkson, convolution, superposition, and

fast superposition algorithms. For each patient, six points are selected to compare dose calculation for different algorithms, which used include isocenter of PTV, 3cm from isocenter, 0.5cm under skin, low and high density, and point under the block. Commercially available CMS XiO (Computerized Medical Systems, USA) Planning system was used for planning purposes.

## **Dose Calculation Algorithms**

Four calculation algorithms used to calculate the radiation dose for all plans which were formed through this study. The XiO's fast-Fourier transform (FFT) convolution algorithm, and the superposition (Wiesmeyer and Miften)<sup>5</sup>. Algorithms are comparable that, they both calculate the dose by considering the total energy released in the patient with Monte Carlo-generated energy deposition kernels, computed by Mackie et al.<sup>6</sup>. The kernel is the dose matrix created per unit TERMA at the collaborating sites. Total energy released per unit mass (TERMA) produce from the mass attenuation coefficient and the essential energy flounce<sup>7</sup>. The selection of dose calculation algorithms is a serious respect when using "highended" planning procedures and comparing one method with another<sup>8-10</sup>.

## **Clarkson Sector Integration Algorithm**

One of the most fundamental modified algorithm uses patient data, treatment machine data, and setup information to simulate dose distributions inside the patient. The patient data contain relative electron density information which characterizes a sector of the patient. These have been earlier created by assignment density values in areas incorporated by marking contours or by granting the CT to comparative electron density conversion of patient image data. For a 3-D plan, only transverse patient data are used in the dose computation; Data of The treatment machine required for the Clarkson algorithm must include a set on the diagonal off-center ratios (OCDs), a band of tissue maximum ratios (TXRs) which comes from TAR or modified TPR values, information about the various beam modifiers and different machines (energy) specific constants. If the beam has a multileaf collimator or customized port; the transmissions and outlines of the blocked areas need to be implanted The algorithm takes into account initial dose correction for inhomogeneity in the patient and the transmission by the wedge and bolus, compensators and blocks. Furthermore, the Clarkson algorithm does not take into account scatter modifications due to differences in field intensity (wedges), patient density, or surface curvature. It contracts into account scatter modifications due to field shape. The Clarkson scatter calculation does not accurately the model dose for forked structures because it does not consider the scatter reduction of the air space between the separated contours. Each ray is calculated individually and then the beams are summed together. The full dose of all beams represent the dose that received by the patient<sup>1</sup>.

## **Convolution Algorithm**

The energy deposited kernels of Mackie et al [5]. Must be interpolated from spherical to Cartesian coordinates on a common grid with the TERMA to complete the FFT convolution. Sampling and interpolation of kernels from spherical to Cartesian coordinates are complicated by steep kernel gradients. Adaptive quadrature techniques certify that the correct energy in and near the interaction point is characterized in the Cartesian coordinates.

Evaluation of calculation outcomes specifies that incorrect doses are found if the effect of spread from the neighbors is omitted over a large enough area. It is essential that data of the patient be denoted over a 3D volume since the scatter considered as a point is based on the 3D volume of the scattering intermediate. The essential volume over which scatter of kernel contributions must be necessitated and the maximum volume used in XiO planning system is close to 30cm in the forward direction, 5cm in the backward direction, and replicate the field size dimension laterally (basically the contributions from all interaction points must be collected). Sharpe and Battista<sup>11</sup> have reported the same ranges, as well as Mackie *et al.*<sup>12,13</sup> have described the same obligatory lateral range.

As well as dose contributions above such a large area needs a significant time for computation. This computation time can be reduced by accomplishment separate calculations; one with the main core for which the computation is done at a high resolution, but over a small region, the other with a scatter kernel where a computation is achieved at a lower resolution but over a large area, as suggested by Mackie *et al.*<sup>5,8</sup>. This method is possible for the primary kernels have particularly large gradients close to the point of interaction, but they make no contribution outside a few centimeters from the interaction point, whereas, the scatter kernels have smaller gradients but contribute the dose over a much larger range. The XiO planning system completes a separate high and low resolution FFT calculated for the primary and scatter kernels, saving a time of about 65% over accomplishment a single calculation at high resolution.

## Superposition Algorithm

The XiO superposition dose accumulated method is a modification of the "collapsed cone" dose calculation process9. As with FFT Convolution, all calculations of superposition are done with coordinates of the beam. The dose in the beam coordinates is interpolated to the user quantified calculation volume. It is conceivable for superposition algorithms to directly imitate the kernel calculation method; that is, to compute deposited energy by dispersion the energy released (TERMA) at the collaboration points to points in the interest volume, according to the distribution obscure by the kernel. This method is recognized as the "interaction point of view". Dissimilar to the FFT convolution algorithms, the superposition algorithm energy deposition kernels can be adjusted to account for fluctuations in electron density. The density scaling method, based on O'Connor's theorem<sup>14</sup> is applied to distort the kernels by finding the average density along the straight-line path between the interaction site and dose deposition site<sup>15</sup>. A good approximation for scattered photons is Density scaling, because the photons move in straight lines and the mass attenuation coefficient scales linearly with the material density (assuming that the atomic number remains unchanged).

## Fast Superposition Algorithm

Spherical kernels, or "dose spread arrays", are cylindrically symmetric and welldefined in terms of rays drawn along zenith and azimuth angles. The spherical kernel computation has been enlarged with the ability to combine (select and sum) adjacent zenith rays in the center. Therefore, it is possible to determine the quantity and direction of zenith rays for the function of adjusting speed/accuracy.

Tradeoffs: The less the rays, the quicker

and less accurate the calculation, The more the rays, the denser and more accurate the calculation. Although the azimuth angles must be evenly spaced, Control of both the number and direction of zenith rays and azimuth rays is possible,. The fast mode provides a fast superposition dose calculation with a speedup factor of 2.5cm at the monetary value of a small loss in accuracy, compared to the "standard" superposition calculation.

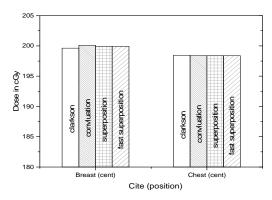
## **Dose Reporting and Evaluation**

All patients plan calculated to receive 200 cGy through two or more treatment field. Weight values ranging from 1 to 100. The weight applied to decrease or increase the dose over the whole volume of the structure. For each patient, dose volume histograms (DVHs) were produced utilizing the CMS XiO planning system for 3DCRT plan. Single dose-volume point were also registered. Minemum dose, maximum dose, and mean dose were estimated for the PTV and OARs. And Uniformity Index Conformity Index were calculated for PTV in all instances. Relative dose volume differences (percentage) between the outcomes of the different dose calculation algorithms were computed. Maximum percent variation between algorithms were recorded for PTV, for all cases. All the sets of treatment plans were assessed utilizing a lot of evaluation parameters, which complied with the evaluation criteria recommended by the international committee on radiation units and measurements (ICRU) Report 62<sup>15,16</sup>. The evaluation parameters included the homogeneity index (HI) and the conformity index (CI). The CI was defined as the quotient of the treated volume and the intensity of the PTV<sup>16</sup>. The conventionally used homogeneity index (H-index) is defined as the ratio of the maximum dose in the PTV to the prescribed dose with a value closer to 1 indicating better homogeneity<sup>17</sup>. The H-index commonly varies from 1 to 1.5 in the practical patient treatment plans. The index's simplicity has led to its being extensively used for quantifying dose homogeneity in tumor volumes. For the evaluation of doses to the OARs, the mean dose was used.

#### **RESULTS AND DISCUSSION**

Fig. (1) Showed insignificant variation in the dosmetric study of the central point of PTV in

both breast and chest cases between four algorithms was used. The maximum percentage of variation was 0. 24% with breast cases, this ratio decreases to 0.03% in cases of chest. As shown in Fig (2) we detect a small variation in dose appeared



**Fig. 1.** Dose in cGy at the center of PTV versus algorithms for breast and chest cases

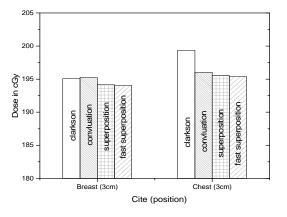
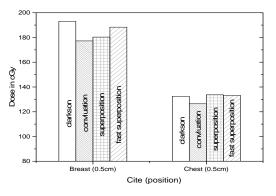


Fig. 2. Dose in cGy at a point far 3cm from center of PTV versus algorithms for breast and chest cases



**Fig. 3.** Dose in cGy at depth 0.5cm from the skin surface versus algorithms for breast and chest cases

at a point far 3cm from the central point which have a percentage 0.6% of breast cases. This variation percentage increase to 2% in the cases of chest. Fig. (3) Declared a large change in the doses which calculated at the point have 0.5 cm depth of skin service with breast cases, where the

205 200 195 Dose in oGy 190 fast superposition superposition fast superposition convluation clarkson superposition convluation 185 clarkson 180 175 Breast (High denisity) Chest (High denisity) Cite (position)

**Fig. 4.** Dose in cGy at point in high density regions versus algorithms for breast and chest cases

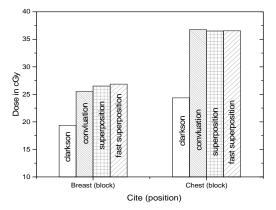
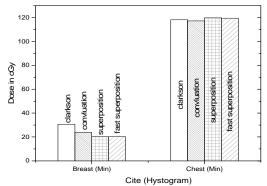
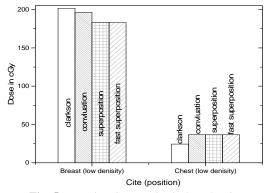


Fig. 6. Dose in cGy at a point under block versus algorithms for breast and chest cases



**Fig. 8.** Minimum dose in cGy calculated from histogram versus algorithms for breast and chest cases

mean percentage of variation in the dose calculation between four algorithms up to 7.9%. At the same point, a significant change was detected in the cases of chest with a dose variation percentage 3.2%, it's clear the variety between four algorithms in same point dose, thus due to the error, in the



**Fig. 5.** Dose in cGy at point in low density regions versus algorithms for breast and chest cases

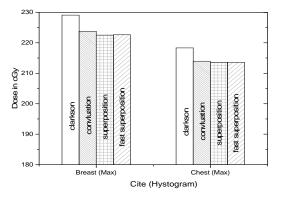


Fig. 7. Maximum dose in cGy calculated from histogram versus algorithms for breast and chest case.

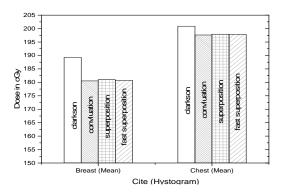


Fig. 9. Mean dose in cGy calculated from histogram versus algorithms for breast and chest cases

dose calculation in the area near skin service with tangential field. We can observe the relative highest in dose which calculate by Clarkson algorithm, where its take in a consider scattering dose from one slid not from hall volume, while this variation decreases to 3.62% in the cases of chest because the opposite field solve partly the error in dose skin calculation. Fig. (4) And fig. (5) represent dose in high and low density regions respectively, A significant variations found with inhomogeneity regions represented by points in high density (bone), low density (lung) where the percentage was 3.01% and 6.2%, respectively, in the cases of chest. On the other hand, in the case of breast these percentage increase to 9.4% at low density region this is due to most of low density area (lung) in the breast field located in penumbra region. While this percentage decreases to 2.62% high density region (bone). As clear in fig (6) At a point in a

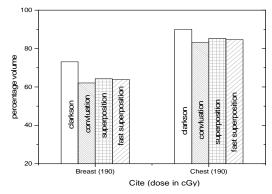


Fig. 10. Percentage of volume, which received dose 200 cGy calculated from histogram versus algorithms for breast and chest cases

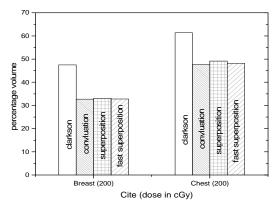


Fig. 11. Percentage of volume, which received dose 200 cGy calculated from histogram versus algorithms for breast and chest cases

protected area in the case of chest, we found a consider percentage of variation about 6.2, the main reason of this variation bake to the dose which calculated by Clarkson algorithm, where it didn't take in consider the internal scatter, we note the main variation with doses calculated by Clarkson algorithm and the other three algorithms have a close value. Figs. (7), (8) and (9) showed Maximum, minimum and the mean dose of PTV for both breast and chest cases which obtained from histogram calculated by each algorithm, the percentage of variation in maximum dose going to 3.26 % and 2.4%, this percentage became 5.2% and 1.2% with a minimum dose of cases of breast and chest respectively. Same variation detected in the mean dose where it founded about 4.4% in the case of breast and 1.63% in the case of chest. The increasing of these parameters with breast cases, related to the PTV of the breast has more inhomogeneity than chest cases. Also histogram gives us a percentage of volume, which received 190cGy (95% of the dose), The difference of these percentages founded as 5.51%, 3.44% of the breast and chest cases respectively as shown in fig. (10). Fig. (11) explained Same behavior founded with 200cGy (100% of the dose) where it appears as 7.4% and 6.8% in the breast and chest cases respectively.

## CONCLUSION

Dose calculated four 28 breast and chest cases, by four different algorithms understudies gives the same values approximately at center point. Except in very small variation with the case of breast, because no difference in field intensity or tissue inhomogeneity at the center in both cases. When we are far from the center a considerable percentage of variation appears, as we see in points, which located at three centimeters from the center. This variation was clear with the doses which calculated by Clarkson algorithm due to the Clarkson algorithm does not take into account scatter modifications due to the field intensity (wedges) so this variation increase in the case of chest. This variation decreases in the case of breast. At the point of depth of 0.5 cm of the skin surface, large variation appear in dose which calculated by convolution algorithm and superposition algorithm recorded in the case of breast this Similarity due to both algorithms compute the dose by convolving the total energy released in the patient so they need a large volume of scatter that's Explains why The variation decrease in the case of chest. The largest value of the dose variation recorded in the case of breast at the point of low density, where the dose calculated by Clarkson and convolution algorithms were convergent, and more than which one calculated by superposition and fast superposition algorithms, which also close.

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