Pediatric Phantom Dosimetry of Kodak 9000 Cone-beam Computed Tomography

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Abstract

Purpose: The purpose of the study was to evaluate the radiation dose of the Kodak 9000 CBCT to different anatomical areas using a pediatric phantom.

Methods: Absorbed doses resulting from maxillary and mandibular region 3x5cm CBCT volumes of an anthropomorphic 10-year-old child phantom (CIRS Inc., Norfolk, VA) were acquired using Optical Stimulated Dosimetry (Nanodot, Landauer Inc., Glenwood, IL) following previously validated protocols. Equivalent doses were calculated for radiosensitive tissues in the head and neck area and effective dose for maxillary and mandibular examinations were calculated following the 2007 recommendations of the International Commission on Radiological Protection (ICRP).

Results: Of the mandibular scans, salivary glands had the highest equivalent dose (1598 µSv) followed by oral mucosa (1263 µSv) then extrathoracic airway (pharynx, larynx and trachea) (859 µSv) and thyroid gland (578 µSv). For the maxilla, salivary glands had the highest equivalent dose (1847 µSv) followed closely by oral mucosa (1673 µSv) then extrathoracic airway (pharynx, larynx and trachea) (1011 µSv) and lens of eye (202 µSv).

Conclusion: Compared to previous research of the Kodak 9000 completed with the adult phantom, the child receives 1-3 times more radiation for mandibular scans and 2-10 times more radiation for maxillary scans.
Introduction

There are several documented uses for cone-beam computed tomography (CBCT) in the pediatric dental field. Cone-beam computed tomography has been utilized to localize developing dentition, visualize resorption in relation to an unerupted tooth, and determine severity of facial trauma. Cone-beam computed tomography has also aided in surgical applications of bony pathoses. Thirty-three percent of 313 cases in pediatric dental patients were for localization of teeth, 19 percent for presence of root resorption, 11 percent for bony pathoses visualization, and finally four percent were facial trauma patients. [1] In a recent retrospective publication, Isman et al. investigated the most common reasons of 329 CBCT in children. They found that dentomaxillofacial anomalies followed by localization of impacted teeth were the most common indications for a CBCT. [2] Dentists and physicians can also benefit from CBCT to visualize the extent of a cleft palate, craniofacial morphology and abnormalities, as well as airway analysis needed for sedation cases. [3] Studies published with regard to pediatric usage of the CBCT mention that in the pediatric population, a smaller field of view (FOV) can satisfy the needs of the prescribing physician or dentist. The smaller the FOV used the less effective dose the patient receives.[3]

Previous research suggest that children are more radiosensitive compared to adults while undergoing dental radiography. Ludlow et al. report effective doses 36% greater in children compared to adults when undergoing a CBCT. [4] They show average effective doses for the maxilla at 53 µSv and average effective doses for the mandible at 102 µSv for an adult phantom. For a child phantom, average effective doses for the maxilla were 67 µSv and average effective doses for the mandible were 128 µSv. [4] Dosages of common dental radiographs, including bitewings and panoramic radiographs, range from 1-20 µSv and 4-30 µSv, respectively. [5]
Therefore, a patient receives a larger amount of radiation while undergoing a CBCT compared to other dental radiographs. Several studies have been published in the area of dosimetry using CBCT with an adult phantom, but there is a lack of publications with pediatric phantoms.

The purpose of the study was to evaluate the radiation dose of the Kodak 9000 CBCT to different anatomical areas using a pediatric phantom with the hypothesis that the child will receive more radiation compared with previous similar studies using an adult phantom.
Methods

Dosimetry is best expressed in terms of “tissue equivalent dose” and “total effective dose”. Tissue equivalent dose \( (H_T) \) is the absorbed dose of the tissue adjusted for the radiation weighting factor. It is calculated by the product of absorbed dose \( (D_T) \) and the radiation weighting factor \( (W_R) \) and expressed in millisieverts or microsieverts. Total effective dose is the calculation the International Commission on Radiological Protection (ICRP) chooses to use to compare differing exposures. [4] It is calculated by taking the sum of the products of the tissue weighting factor \( (W_T) \) and the tissue equivalent dose \( (H_T) \). According to Ludlow et al., this calculation reflects the most radiosensitive tissues and their weighting factor expressing a degree of sensitivity for each tissue and is commonly expressed in millisieverts or microsieverts. [4] The higher the weighting factor, the more radiosensitive the organ is.

A device used for evaluating dose due to exposure from ionizing radiation during dental radiographic examination is an imaging phantom. For this study, an anthropomorphic head and neck phantom (ATOM Max, CIRS, Inc, Norfolk, VA, USA) simulating the approximate size, body type, and mass of an average ten year old child was used to acquire dosimetry data (Figure 1). The phantom contains materials of varying densities which provide attenuation characteristics representative of the varying human tissues, glands, and organs located within the head and neck. The phantom is sectioned into axially oriented slabs (25 mm thick), which permits access to specific tissues and anatomical locations of interest (Table 1). Slabs are modified to accept dosimeters at each of the internal and external sites. During the imaging process, the phantom was oriented so that the sectioned planes were parallel to the floor.
Dosimetry was recorded using optically stimulated luminescence (OSL) dosimeters (Nanodot, Landauer, Glenwood, IL, USA). OSL dosimeters respond to ionizing radiation by storing energy in proportion to the amount of x-ray energy to which they are exposed. Each dosimeter is encased in a light-tight plastic holder measuring approximately 1mm x 10mm x 10mm. This case prevents any ambient lighting from reaching the dosimeter and therefore causing skewed data. Sets of 24 dosimeters, each corresponding to a specific organ or tissue of interest, were grouped and coded for identification. Each set was cleared of stored energy using a light source (LED light pad) for at least 24 hours prior to establishing baseline readings. Seven dosimeter sets were used during the study, one served as a control set.

The Kodak 9000 (Carestream Dental LLC, Atlanta, GA, USA) has one FOV, 50mm x 37mm. The voxel size used was 0.076 mm. This voxel size will provide better image resolution and detail compared with a bigger voxel size (0.4). However, the radiation with smaller voxel sizes is higher. For maxillary techniques, 12 scans were completed using the same dosimeter set with the FOV focused on the permanent maxillary left first molar (#14). This procedure was repeated two more times, each time utilizing a different set of dosimeters with the same FOV location. Each dosimeter set was averaged to calculate the dose per examination. The same technique was used for mandibular exposures with the FOV focused on the permanent mandibular left first molar (#19). Since the Kodak 9000 has a smaller FOV compared to other CBCT units, more exposures were completed. “Smaller FOVs require more exposure repetitions because more dosimeters are outside of the field of direct exposure and absorb only small quantities of scatter radiation.” [4] All scans were acquired using the same “child” setting set by the manufacturer: 75 kV and 8 mA.

Dosimeters were read with a portable reader (MicroStarii, Landauer, Glenwood, IL, USA). The reader was calibrated initially with a set of dosimeters, supplied by the manufacturer, which
had been exposed to known amounts of energy. Reader performance was checked before each use. Average and standard deviation of each set of dosimeters were calculated. Effective dose (\(\mu\text{Sv}\)) was calculated by using the same methodology published by Johnson B et al. and applying 2007 ICRP tissue weighting factors. [6]
Results

Table 2 represents the tissue equivalent doses and the total effective dose for each scan, three for mandibular scans and three for maxillary scans. Table 3 represents the average and standard deviation for tissue equivalent doses and total effective dose for mandibular and maxillary scans. The average effective dose of the mandibular scans was 65.4 µSv ± 3.2 µSv. The average effective dose of the maxillary scans was 53.2 µSv + 2.5 µSv. Graph 1 shows the average equivalent doses of tissues for both mandibular and maxillary scans. Of the mandibular scans, the largest equivalent dose per organ was seen in the salivary glands (parotid, submandibular, and sublingual) (1598.5 µSv ± 107.9 µSv) followed by oral mucosa (1263.3 µSv ± 104.3 µSv), extrathoracic airway (pharynx, larynx and trachea) (859.4 µSv ± 55.1 µSv), and the thyroid gland (578.9 µSv ± 73.4 µSv). Of the maxillary scans, the largest equivalent dose per organ was seen in the salivary glands (parotid, submandibular, and sublingual) (1847.8 µSv ± 61.4 µSv) followed by oral mucosa (1673.0 µSv ± 82.7 µSv), extrathoracic airway (pharynx, larynx and trachea) (1011.4 µSv ± 45.4 µSv), and the lens of the eye (202.5 µSv ± 16.1 µSv).
Discussion

Stochastic effects of radiation, or the damage to the DNA causing cancer or other heritable defects, are an adverse outcome based on the frequency of radiation. [5] The larger the equivalent dose to a tissue, the more likely stochastic effects occur. However, for head and neck radiographs such as the CBCT where effective dose is less than 0.1 mSv (100 µSv), the risk of stochastic effects are negligible. [7] It is important to note that effective dose of this study does not correlate to a specific patient, but more to a “reference patient” of an average 10 year old child, as there are known difference with age and sex. [7]

Pauwels et al completed a study in 2012 using the adult phantom testing numerous CBCT machines, including the Kodak 9000. The FOV of the Kodak 9000 specifically focused on the lower jaw molar region resulted in an effective dose of 40 µSv and an equivalent dose to the salivary glands 709 µSv. [8] Compared to this study, the child phantom with the FOV focused in the same location resulted in 1.6 times greater effective dose and 2.3 times greater equivalent dose to the salivary glands.

In a meta-analysis completed by Ludlow et al., numerous CBCT machines were analyzed based on FOV size and default or standard settings based on manufacture of the machine, some utilizing the adult and child phantom. The Kodak 9000 (CS 9000) machine was analyzed only using the adult phantom with standard adult settings for both maxillary and mandibular scans. Of those findings, the maxillary effective dose ranged from 5-19 µSv and the mandibular effective dose ranged from 22-40 µSv. [9] These reported effective doses, when compared to this study of
the child phantom, reveal that the child receives 2.8-10 times and 1.6-2.9 times greater for the maxillary and mandibular scans, respectively. Therefore, the child receives roughly 2-10 times more radiation overall when undergoing a scan of the maxilla and 1-3 times more when undergoing a scan of the mandible compared to an adult.

Salivary glands were also the organ to receive the largest equivalent dose of the adult phantom based on the meta-analysis of the Ludlow et al. [10] The salivary glands specifically received 130-523 µSv with scans of the maxilla and 633-1037 µSv with scans of the mandible. This is 3.5-14.2 times more radiation to the salivary glands of a child undergoing a maxillary scan and 1.5-2.5 times more radiation to the salivary glands of a child undergoing a mandibular scan.

The salivary glands were not incorporated into the ICRP calculation of effective dose until 2007. The 2007 ICRP Guidelines include salivary glands and updated tissue-weighting factors for other organs. [11] Review of dosimetry literature prior to 2007 shows lower effective doses for both pediatric and adult phantoms. Ludlow et al found an increased effective dose of 32-422 percent with the use of the 2007 ICRP guidelines compared to the previous guidelines. [12]

In order to better understand how much radiation a child is exposed to while having a CBCT with the Kodak 9000, effective doses can be compared to the effective doses of common intraoral radiographs (posterior bitewings). Johnson B et al. calculated the effective dose (µSv) for a 12 yer old child using a F-speed film and with rectangular collimator in 5 µSv. [6] We found that a 12 year old child receives an average effective dose 65 µSv with a CBCT limited to the mandible. Ten times more effective dose when undergoing a CBCT compared to bitewings with rectangular collimation.

Further work needs to be completed in the field of child phantom dosimetry with other CBCT machines. Due to the differing manufacturer settings of CBCT machines and variable
scanning options of CBCT machines, more research is required to fully understand the amounts of radiation a child is exposed to. This study is limited to one CBCT machine with one FOV option. Future dosimetry research can be completed using other machines that have been studied with the adult phantom for additional comparisons to be made.

Conclusion

Cone-beam computed tomography should be used judiciously in pediatric patients due to the overall amount of radiation exposure.

1. Pediatric patients receive up to 10 times more radiation when compared to adult patients undergoing a CBCT with the Kodak 9000.

2. Pediatric patients receive the most radiation to the salivary glands with both maxillary and mandibular scans with the Kodak 9000.

Acknowledgements

The research leading to these results was possible with use of the pediatric phantom, OSL dosimeters, and the dosimeter reader loaned from the North Carolina Oral Health Institute in Chapel Hill, North Carolina.
Table 1

<table>
<thead>
<tr>
<th>OSL ID</th>
<th>Child Phantom Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Calvarium anterior (2)</td>
</tr>
<tr>
<td>2</td>
<td>Calvarium left (2)</td>
</tr>
<tr>
<td>3</td>
<td>Calvarium posterior (2)</td>
</tr>
<tr>
<td>4</td>
<td>Mid brain (2)</td>
</tr>
<tr>
<td>5</td>
<td>Mid brain (3)</td>
</tr>
<tr>
<td>6</td>
<td>Pituitary (4)</td>
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<td>7</td>
<td>Right orbit (4)</td>
</tr>
<tr>
<td>8</td>
<td>Right lens of eye (4-5)</td>
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<tr>
<td>9</td>
<td>Left lens of eye (4-5)</td>
</tr>
<tr>
<td>10</td>
<td>Right maxillary sinus (5)</td>
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<tr>
<td>11</td>
<td>Left nasal airway (5)</td>
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<td>12</td>
<td>Right parotid (6)</td>
</tr>
<tr>
<td>13</td>
<td>Left parotid (6)</td>
</tr>
<tr>
<td>14</td>
<td>Left back of neck (6)</td>
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<td>16</td>
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<td>17</td>
<td>Right submandibular gland (7)</td>
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<tr>
<td>18</td>
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<td>Center sublingual gland (7)</td>
</tr>
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<td>Center C spine (8)</td>
</tr>
<tr>
<td>21</td>
<td>Thyroid superior left (8)</td>
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<tr>
<td>22</td>
<td>Thyroid left (9)</td>
</tr>
<tr>
<td>23</td>
<td>Thyroid right (9)</td>
</tr>
<tr>
<td>24</td>
<td>Esophagus (9)</td>
</tr>
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</table>

Location of dosimeters inside the pediatric phantom [axial slice indicated by ( )]
Table 2

<table>
<thead>
<tr>
<th>Exam/Location (μSv)</th>
<th>Mandible 1</th>
<th>Mandible 2</th>
<th>Mandible 3</th>
<th>Maxilla 1</th>
<th>Maxilla 2</th>
<th>Maxilla 3</th>
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<tbody>
<tr>
<td>Bone Marrow</td>
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<td>28.3</td>
<td>19.5</td>
<td>15.1</td>
<td>20.6</td>
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<tr>
<td>Thyroid</td>
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<td>661.3</td>
<td>550.0</td>
<td>145.1</td>
<td>128.8</td>
<td>101.6</td>
</tr>
<tr>
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<td>15.4</td>
<td>20.4</td>
<td>16.4</td>
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<td>6.7</td>
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<td>Skin</td>
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<td>31.3</td>
<td>31.3</td>
<td>31.9</td>
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<td>Bone surface</td>
<td>133.7</td>
<td>152.8</td>
<td>131.5</td>
<td>88.7</td>
<td>68.7</td>
<td>94.7</td>
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<td>Salivary glands</td>
<td>1569.0</td>
<td>1515.8</td>
<td>1720.5</td>
<td>1889.8</td>
<td>1777.4</td>
<td>1876.3</td>
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<tr>
<td>Remainder</td>
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<td>162.2</td>
<td>184.8</td>
<td>221.7</td>
<td>202.5</td>
<td>217.8</td>
</tr>
<tr>
<td>Brain</td>
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<td>12.1</td>
<td>16.4</td>
<td>38.1</td>
<td>36.5</td>
<td>37.2</td>
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<tr>
<td>Lymphatic nodes</td>
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<td>45.1</td>
<td>48.5</td>
<td>50.9</td>
<td>46.3</td>
<td>49.6</td>
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<td>823.9</td>
<td>831.5</td>
<td>922.9</td>
<td>1050.0</td>
<td>961.4</td>
<td>1022.7</td>
</tr>
<tr>
<td>Muscle</td>
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<td>45.1</td>
<td>48.5</td>
<td>50.9</td>
<td>46.3</td>
<td>49.6</td>
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<td>Lens of eyes</td>
<td>54.2</td>
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<td>Pituitary</td>
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<td>29.6</td>
<td>71.0</td>
<td>68.5</td>
<td>67.8</td>
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<tr>
<td>Effective Dose</td>
<td>61.8</td>
<td>67.5</td>
<td>67.0</td>
<td>55.5</td>
<td>50.6</td>
<td>53.3</td>
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</table>

Tissue equivalent doses and effective dose for standard parameters of Kodak 9000
Table 3

<table>
<thead>
<tr>
<th>Exam/Location (µSv)</th>
<th>Mandible Average</th>
<th>Mandible Standard Deviation</th>
<th>Maxilla Average</th>
<th>Maxilla Standard Deviation</th>
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</thead>
<tbody>
<tr>
<td>Bone Marrow</td>
<td>29.8</td>
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<td>Esophagus</td>
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<td>0.4</td>
</tr>
<tr>
<td>Skin</td>
<td>3.8</td>
<td>0.6</td>
<td>31.2</td>
<td>0.8</td>
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<tr>
<td>Bone surface</td>
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<td>11.7</td>
<td>84.1</td>
<td>13.6</td>
</tr>
<tr>
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<td>13.8</td>
<td>2.3</td>
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<td>Lymphatic nodes</td>
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<td>48.9</td>
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<td>48.9</td>
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<td>Oral mucosa</td>
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<td>82.7</td>
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<td>3.2</td>
<td>53.2</td>
<td>2.5</td>
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Average and standard deviation for tissue equivalent doses and effective dose for standard parameters of Kodak 9000
References