

Full Length Research Paper

Environmental assessment, instrumentation-quality tests of radiological equipment and human health implications

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The health implications of human exposures to ionizing radiation may be deleterious on the immunity mechanism, particularly when large but sub-lethal doses are applied over a period of time. Quality control tests of X-ray equipment and environmental monitoring of a private radiological center (Moroko X-ray Center, Lagos) was carried out. The results showed that radiological personnel and the facilities for safety were grossly inadequate. Dose rates of 3.0 to 3.6 μSv were recorded at the entrance door when the door is closed and opened respectively. The dose rates at the adjacent room and the waiting lobby were at least a factor of 60 higher than the background dose rate each, indicating higher risk to visitors and personnel at the center. Investigation of the equipment using a calibrated multi-purpose non-invasive X-ray test device, Victoreen model 4000M+ and radiation monitor device minirad model 1000+, showed lower values against recommended limits of the International Commission of Radiological Protection (ICRP). However, general overhauling of the facilities was recommended as the X-ray equipment is older than the recommended age by the Nigerian Nuclear Regulatory Authority (NNRA). A follow-up study indicates improved facilities and safety measures.

Key words: Quality control, diagnostic facilities, radiology, health, immunity

INTRODUCTION

Quality control (QC) in atomic radiation comprises the regular tests that must be carried out on major components of the radiation machine to ensure its optimum performance within the system as a whole (West, 1993; CRCPD, 2001; AAPM, 2002). Constant monitoring and routine Quality Control procedures and recommend tests of pieces of diagnostic Imaging equipment are a vital requirement (Cancer Care, 2004).

X-rays are a form of ionizing radiation capable of traveling through materials. Since the discovery by German Physicist Wilhelm Conrad Roentgen in 1895, X-ray technology has been an invaluable tool in medicine, industry, agriculture, scientific research, security and safety. For instance, it can be used to observe broken bones and swallowed objects (Zuur, 2002), track blood flow in patients (Taber et al., 2005), and check for

cavities. X-rays can also be used to diagnose cancer, kill bacteria in food (Grolichová et al., 2004), analyze the structure of crystals or distant stars (LaRouche, 2010; Martinson, 2010), and scan baggage at airports (Zhumadilov et al., 2008).

X-ray production, detection, image processing, and image viewing are some of the major systems in diagnostic radiology to which major quality control can be applied. These tests must be coupled with routine monitoring of final image quality and the environment (BIR, 1988). The purpose of a radiation monitoring programme is to identify all sources of radiation exposure within an operation area, to assess the level of radiation exposure of the employee and members of the public (UNSCEAR, 2005), for the timely detection of changes in radiation parameters which may lead to increased exposures, and to produce sufficient information for optimization purpose (Zoetelief et al., 2006). Equipment quality control unit carries out evaluation of equipment performance to ensure proper image quality, as well as patient and operators' safety.

In spite of the fact that X-rays like other ionizing radiation have beneficial usage in various fields, improper handling and lack of monitoring system may give rise to undesirable effects, some of which have to be weighed against the benefits in application. The energy associated with electromagnetic waves upon interaction with exposed matter may generate heat which can damage cell material (Zuur, 2002). Furthermore, interactions between radiation and matter can result in the production of free radicals which can interfere with the DNA present in cells (Zuur, 2002). The damage done is sometimes irreparable. However, oftentimes the damage is irreparable or an incorrect repair can occur. DNA damage due to relatively low doses may cause so called 'stochastic effects' which can be either tumour induction or genetic effects (Zuur, 2002; Berrington et al., 2004; UNSCEAR, 2006). It has been observed that following radiation injuries, the normal microbe flora of the gut, skin and respiratory tract are disconcerted (Leone, 1962). Exposure to sublethal doses of radiation caused the bacterial inhabitants of the gastrointestinal tract to invade the tissues and organs and set up bacteremia which may lead to death (Brown, 1962) or of serious morbid effect in the exposed individual. Genetic alterations due to energy deposited in irradiated cells can arise in non-irradiated cells by so-called non-targeted effects (Morgan, 2003) by receiving signals produced by irradiated cells or the descendants of irradiated cells (Coates et al., 2008; Wright, 2010). In addition, exposure to radiation may lead to resistance in terms of combating pathogenic microorganisms. The mechanism involved in such instance is that of the deleterious effect of ionizing radiations on the immune system. The immune system which is composed of lymphocytes and other accessory cells which are all over the body function to recognize antigens (foreign bodies), neutralize or quench their

activities thereby preventing infections and tumours. Lymphocytes are radiosensitive and exposure to low doses and low rates of ionizing radiation might lead to immunosuppression (UNSCEAR, 2006), hence reduced defense-ability.

MATERIALS AND METHODS

In X-ray production, some of the variables investigated during quality control include; peak tube voltage (kVp), product of tube current and exposure time (mAs), beam filtration, automatic exposure devices (AEDs), machine output, X-ray beam/alignment and focal spot. These variables are initially checked to establish a baseline for Quality Assurance (QA) programme (AAPM, 2002). Thereafter, regular testing is contrived to AEDs machine output and beam alignment. Certain test is expected to be carried out on weekly basis (AAPM, 1991) on certain devices such as AED since it has the tendency to lose calibration over a period of time and this will affect both image quality and patient dose. Another variable that requires weekly measurement is the radiographic output.

There are over 4000 x-ray machines in the Nigeria with less than 5% of them under regulatory control, thereby posing serious challenges (Elegba, 2006) which in turn affect the quality of patient dose and image quality.

In this study we investigate the efficiency/output of X-ray machine in a private radiology centre in Nigeria. The intention is to compare the degree of conformity of current practice in the center to requirements of the International Basic Safety Standards [BSS] and hence, guide against negative consequential effects of non-compliance on human health.

Before measurements were carried out the devices to be used were standardized by calibration at the Secondary Standard dosimetry Laboratory (SSDL) of the Nigerian Institute of Radiation Protection and Research (NIRPR).

A multi-purpose measuring non-invasive X-ray test device, Victoreen model 4000M+ was used in the collection of data that bothered on the Peak Tube Voltage Kvp, Timer (Accuracy and Reproducibility), Consistency of output and Linearity of X-ray output. The tests carried out on the X-ray facility were according to the manufacturer's guideline (Victoreen, 2008). The device was employed to determine, the voltage and timer (accuracy and reproducibility), as well as the X-ray machine's current and output dose. The environmental monitoring was carried out using calibrated radiation monitor device minirad model 1000+ GM Survey Meter (Thermo scientific, 2008).

RESULTS

General observations

The observations made as shown in Table 1 is a reflection of the result of radiation dosage at the entrance lead lined door measured which is between 3.0 to 3.6 $\mu\text{Sv/h}$, the dose rate at the adjacent room and the waiting lobby were a factor of 60 higher than the background dose rate. Within the cubicle a dose rate of 2.0 $\mu\text{Sv/h}$ was measured. Hazard warning light and signs were also not provided; Personnel Monitoring data using thermoluminescence dosimeter (TLD) badges were not available. This fall short of the expected standards required by the National and International Regulatory

Table 1. General observations.

S/N	Observations	Centres abbreviation	
		MXC	
		Yes	No
1	Main door to x-ray room (Lead Lined)	X	-
2	Main door to x-ray room (Lead efficient)	-	X
3	Second Door to x-ray room (lead lined)	nil	-
4	Second door to X-ray room lead lined (efficient)	nil	-
5	Cubicle Type Lead Wood	X	-
6	Cubicle Type Lead wall	-	-
7	Cubicle Efficient	-	X
8	Cubicle window (efficient)	-	X
9	Door Interlock provided	-	X
10	Door (close Automatically)	-	X
11	Provision of lead apron	-	X
12	Lead Apron Efficient	-	X
13	Hazard warning light Provided	-	X
14	Hazard warning light functional	-	X
15	Hazard warning sign (Displayed)	-	X
16	Functional Air-Conditional Provided	-	X
17	Personnel Monitoring (TLD badges) available	-	X
18	Qualified Radiographers available	X	-
19	Darkroom and x-ray room Interconnected	-	X
20	Darkroom temperature controlled	-	X
21	Thoroughfare (Prohibited)	-	X
22	Log book available	-	X
23	X-ray machine over 10 years?	-	X
24	Space of x-ray room (adequate)	-	X
25	Collimator light functional	X	-

Y (X) = Yes, N (X) = No, nil = not available.

Table 2. Quality control tests carried out.

S/N	Radiological tests	
1	Peak Tube Voltage Kvp (Accuracy & Reproducibility).	
2	Timer Accuracy and Reproducibility.	
3	Optical-Radiation field	
4	Light/Radiation Beam Alignment	
5	Constancy of output	
6	Linearity	i Current (mA) ii Timer
7	Reproducibility of X-ray of output	

CRCPD (2001).

Bodies (IAEA, 1996; NIBIRR, 2003).

Quality control test

Table 2 shows the quality control (QC) tests performed

on the equipment. The X-ray equipment was found not to measure accurately and does not reproduce set values. At the following kvp settings 65, 70, 75, 80 lower values of 57, 62, 69, 71 respectively were measured. Tube voltage should have 5% accuracy and 2.5% precision (Table 3). The AEC Optical density of the film should be between 1.10 OD and 1.50 OD (van den Berg et al., 1998)

For the time accuracy, none of the reading measured was within $\pm 5\%$ of the settings. The measured kvp values are not linear compared with that of the settings. For linearity test $<0.5\%$ or $<0.43 \mu\text{Gy}$ (0.05 mR) deviation from linearity for any exposure within the readout range of 17.4 μGy (2 mR) to 86.9 Gy (10 R), is at a given exposure rate (AAPM, 1991). The values measured could also not be reproduced.

DISCUSSION

The accuracy and reproducibility of kVp measured values

Table 3. Quality control tests on X-ray tube and generator.

Type of tests	Settings			Measured values		
	Kvp	Time (s)	mAs	Kv efficient	Exposure time (s)	Output dose (mR)
Accuracy (Kvp)	65	0.04	10	56.78	0.0313	34.78
	70	0.04	10	62.33	0.0313	37.57
	75	0.04	10	68.78	0.0313	41.32
	80	0.04	10	71.34	0.0313	44.14
Accuracy (time)	70	0.02	10	61.97	0.0415	18.82
	70	0.04	10	62.64	0.0894	38.56
	70	0.08	10	62.55	0.1409	77.22
	70	0.16	10	63.32	0.1267	119.85
Consistency (Kvp)	70	0.05	10	61.97	0.0415	18.82
	70	0.1	10	62.64	0.0894	38.56
	70	0.15	10	62.55	0.1409	77.22
	70	0.2	10	63.32	0.1267	119.85
Linearity (ma)	70	0.04	10	62.33	0.0313	37.56
	70	0.04	20	63.53	0.0313	76.13
	70	0.04	30	63.32	0.0313	151.62
	70	0.04	40	63.46	0.0313	299.89
Reproducibility (Kvp)	70	0.04	10	63.44	0.0312	37.57
	70	0.04	10	62.61	0.0323	37.57
	70	0.04	10	62.77	0.0350	37.57
Reproducibility (timer)	70	0.04	10	63.44	0.0312	37.57
	70	0.04	10	62.61	0.0323	37.57
	70	0.04	10	62.77	0.0350	37.57

were not within acceptable range as previously stipulated (BIR, 1988; NRPB, 1988). From these results, regular evaluation of the health status of health workers who may be exposed to low radiation doses is warranted. The quality control tests of an X-ray unit are to be undertaken by a Medical Physicist with the purpose of safety and dose optimization (AAPM, 1991). The x-ray tube voltage (Kilo volt [peak]) has a significant effect in the image contrast, the optical density and the patient dose, variations between the stated kilovolt (peak) and the x-ray beam quality must be within $\pm 5\%$. For the x-ray machine investigated, there is deviation of the measured kVp from the set values on the control panel. The kVp measurements from the X-ray machine as shown in Table 3 are $< 5\%$ accuracy which is in discordance to the limiting values as reported by van den Berg et al. (1998). The QC test carried out on the consistency of the machine showed non-compliance. Quality management programme needed for radiation safety policies and procedures were not put in place. Record showed that Radiation Safety Officers and Medical Physicists were

not engaged in the activities of the centre. Medical Physics Experts will contribute to maintaining and improving the quality, safety and cost-effectiveness of healthcare services through patient-oriented activities requiring expert action, involvement or advice regarding the specification, selection, acceptance testing, commissioning, quality assurance/control and optimised clinical use of medical radiological devices and regarding patient risks from associated ionising radiations (AAPM, 1991; NIBIRR, 2003; MPE, 2012). The film exposed revealed light and x-ray field misalignment which should not exceed 2% (CRCPD, 2001); in this case it is above the 2% of the source-to-image distance (SID) of both the length and the width of the film. In the light beam alignment test, the film produce revealed the edges light and X ray beam to be within 1% focus–film distance (van den Berg et al., 1998). The gross deviation from acceptable standards may be an indication of exposure of personnel and visitors to radiation risks.

From the foregoing, it is generally recommended that the center be completely overhauled.

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