



## Effect of Gamma Irradiation on the Shelf Life and Quality of Mutton

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### ABSTRACT

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The present study was undertaken to assay the effects of gamma irradiation on fresh mutton compared with non-irradiated mutton. For this purpose the meat samples were divided into four treatments. The experiment was conducted on fresh mutton treated with <sup>60</sup>Co gamma irradiation having dose of 0 (T<sub>0</sub>), 1.5 (T<sub>1</sub>), 2 (T<sub>2</sub>) and 4 kGy (T<sub>3</sub>) and stored in 0, 30 and 60 days at -20°C to estimate the effect on proximate, sensory, physicochemical, biochemical and microbial changes in mutton. Proximate analysis (DM, CP, EE, and Ash), physicochemical and bio-chemical tests (pH value, cooking loss, PV, TBARS and FFA), sensory tests (color, flavor, tenderness, juiciness, overall acceptability) and microbial test (TVC, TCC and TYMC) were done to evaluate the effect of irradiation on shelf life and quality of mutton during preservation. The mutton samples were stored up to 2 months and evaluated at 0, 30 and 60<sup>th</sup> day. Irradiation increased DM, EE and decreased ash of mutton. CP was decreased with treatment. The PV, TBARS, FFA values and Cooking loss were increased with treatment. But raw pH was decreased with irradiation. There were significant differences in color and flavor with treatment. There were no significant differences in tenderness, juiciness and overall acceptability with treatment. The TVC (logCFU/g), TCC (logCFU/g) and TYMC (logCFU/g) were decreased with treatment. As a result, it can be concluded that irradiation increased shelf life of mutton to maintain nutritional quality of meat. Among the treatments, the highest irradiation dose (4.0 KGy) found the best results in terms of overall acceptability, microorganism control and for shelf life extension of mutton.

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## Introduction

Sheep is the 3<sup>rd</sup> largest ruminant animal in Bangladesh and plays a significant role in meat production in Bangladesh. About 23% of ruminant animals were slaughtered every year and among them 7% of these animals were slaughtered in a single day at Eid ul Adha in Bangladesh (Akter et al., 2009). A surplus amount of meat produced during Eid ul Adha needs to be preserved safely for further consumption. Several methods were used to preserve meat such as drying (Akhter et al., 2009), freezing (Bithi et al., 2020; Modak et al., 2009; Rahman et al., 2014), curing (Inguglia et al., 2018), smoking (Bruneau et al., 2005), and irradiation (Haque et al., 2017; Islam et al., 2019; Rima et al., 2019). Irradiation is a physical means of sterilization or decontamination where products are exposed to gamma rays. Gamma-rays are electromagnetic waves emitted from radioactive isotopes such as cobalt-60 or cesium-137 that have strong penetration (Shin et al., 2014). Electron-beams are mechanically generated by

electron accelerator and directly irradiate products, it has advantages of control convenience and favorable perception consumer acceptance, but weak penetration power compared to gamma-ray. A recent study has indicated that three radiation sources (gamma-ray, electron-beam and X-ray) led to different efficacies on microbial properties as well as color characteristics and lipid oxidation stability. Gamma-ray irradiation increases the redness of pork sausages; on the other hand it decreases by electron-beam irradiation at higher dose level. Electron-beam irradiation effectively reduced the total aerobic bacteria in beef patties whereas, gamma-ray considerably decreased microbes in pork sausages as irradiation level increased (Ham et al., 2017). Several methods which include cooking, freezing, fermenting, salting, drying and pickling (Choi et al., 2009; Kim et al., 2009) have been used to reduce the number of microorganisms and increase the shelf-life and safety of meat. However, one of the most

promising approaches to improve microbial safety of meat would be the use of low or medium-dose (1–10 kGy) irradiation (WHO, 1999). Several researchers have stated that irradiation can control efficiently the growth of pathogenic and spoilage bacteria in meat (Zhu et al., 2005; Islam et al., 2019).

Irradiation extends shelf life, destroy pest, reduces infection from food borne pathogens or parasites and has been shown to be a healthy state (Thayer et al., 1995). Oxidation of lipid and lipid substances are the major causes of reducing the quality of meat during storage. Adoption of irradiation on meat induces the production of free radicals which react with oxygen, leading to the formation of carbonyls that helps to maintain the nutritional and sensorial characteristics of meat. Food irradiation technology has been practically used as an effective preservative method to not only reduce initial microbial contamination, but also delay microorganism growth during storage in meat products (Kim et al., 2012; Roberts, 2014). The U.S. Food Safety and Inspection Service approved irradiation of red meat in 1997 (FSIS, USDA, 2008). Irradiation doses required to kill pathogens can cause undesirable changes in meat color, flavor and odor (Nam et al., 2003). Irradiation is the most effective technology in eliminating these pathogens in meat products. Most of the irradiation studies so far are mainly focused on raw meat and meat products without additives and little information on the quality changes of ready to eat meat products by irradiation are available. Irradiating uncooked meat produced a characteristic blood-spattered and a pleasant aroma that remained after cooking (Hashim et al., 1995).

Lately Bangladesh is producing 72.60 Lakh MT where as demand of 72.14 Lakh MT (DLS, 2018) where mutton is contributing 12% of total meat production. It indicates that Bangladesh is self sufficient in meat production and the amount of surplus is 0.46 Lakh MT. As a result now we have opportunity to seek foreign markets to export our excess meats. It is essential to develop a way to increase the shelf life and quality of raw meat to overcome the international trade barrier. In this case, irradiation can be an effective way to increase the shelf life and safety of meats. At present in Bangladesh, no research has yet been carried out on irradiation of mutton. Therefore, the present study was undertaken to determine the effect of gamma irradiation on sensory, proximate component, biochemical and microbial parameters of mutton and also to find out the suitable level of irradiation dosage on mutton to increase the shelf life.

## Materials and Methods

### Sample Collection, Processing and Irradiation

Boneless mutton from freshly slaughtered three indigenous Bangladeshi sheep of same age, sex and body condition was collected from local market of Bangladesh Agricultural University, Mymensingh. Meat samples (500gm from each sheep) were collected from the same location (shoulder) of all three sheep. Then the samples were immediately transferred to the Laboratory of Animal Science. Fat, ligaments, bone and tendons were trimmed with sharp knives and cleaned with distilled water. After that, the meat samples were divided into four groups and

labeled with specific radiation dose. Each group was exposed to the irradiation dose of 0 (T<sub>0</sub>), 1.5 (T<sub>1</sub>), 2 (T<sub>2</sub>) and 4 kGy (T<sub>3</sub>) at the Bangladesh Institute of Nuclear Agriculture (BINA) using Cobalt 60 GC-5000 (BRIT, India) machine for irradiation source and stored at -20°C for pending analysis.

### Quality Analysis

Different proximate, sensory and bio-chemical measurements were examined. Mutton samples were through panel test consisting of 6-8 members. Panelists were selected among department staff and students and trained according to the American Meat Science Association guidelines (AMSA, 1995). The sensory questionnaires measured on a 5-point balanced semantic scale (weak to strong) for the following attributes color, smell, tenderness, juiciness, and overall acceptability. The judges evaluated the samples based on AMSA guidelines. Evaluation was done in individual booth under controlled conditions of light, temperature and humidity. Before the evaluation process, all the panelists participated in an orientation session to familiarize with the meat judging (color, flavor, juiciness, tenderness, overall acceptability) of mutton using an intensity scale. Sensory qualities of the mutton samples were judged after thawing at before and after cook with a 5-point rating scale. Sensory scores were 5 for excellent, 4 for very good, 3 for good, 2 for fair and 1 for poor (Rahman et al., 2012). All samples were served in the Petri dishes. Sensory evaluation was done at 0 day and repeated at 30 and 60 days; up to the end of the refrigerated storage at -20±1°C.

### Proximate nutrients analysis

Proximate composition such as dry matter (DM), ether extracts (EE), crude protein (CP) and ash were determined according to the methods of AOAC (2005). All the parameters were done in triplicate and the mean values were incorporated.

### Physicochemical and Biochemical analysis

Two physicochemical analyses such as pH value of raw meat and cooking loss; three biochemical analysis such as Free Fatty Acid (FFA), Peroxide Value (POV), Thiobarbituric Acid values (TBARS) were measured.

### Raw pH Measurement

The pH value of raw meat was measured using pH meter from raw meat homogenate. The homogenate was prepared by blending 5 g of meat with 10 ml distilled water.

### Cooking Loss

To determine cooking loss, five (5) g sample was weighed and wrapped in a foil paper and it was heated in a water bath for 30 minutes at 75°C. The temperature of water bath was measured by a digital food grade thermometer (HT-2). It needed approximately 30 minutes to attain a temperature up to 75°C. After that the sample surfaces were dried and weighed to measure cooking loss.

The weight of the samples was weighed before and after the heat treatment. Cooking loss is expressed as the ratio of weight loss over the initial weight, using the follow formula:

$$\text{Cooking loss \%} = \frac{w_1 - w_2}{w_2} \times 100$$

Where, w<sub>1</sub> = meat weight before cooking (g) and w<sub>2</sub> = meat weight after cooking (g).

### Free Fatty Acid (%) Analysis

The free fatty acid value of fresh mutton was determined according to the method adopted by Rukunudin et al. (1998) and expressed as percentage. The determination of FFA using the following equation:

$$\text{FFA (\%)} = \frac{\text{Titration volume} \times \text{normality of KOH} \times 28.2}{\text{g of sample}}$$

### Peroxide Value (POV) Analysis (meq/kg)

The Peroxide value (POV) was estimated according to the method adopted by Sallam et al. (2004). The determination of peroxide value was determined using following equation:

$$\text{POV (meq/kg)} = \frac{S \times N}{W} \times 100$$

Where S is the volume of titration (ml), N the normality of sodium thiosulfate solution (n = 0.01) and W the sample weight (g).

### Thiobarbituric Acid Values (TBARS)

Thiobarbituric Acid Values (TBARS) were determined to assess the lipid oxidation status of a supplied sample. It was done in triplicate using the 2-thiobarbituric acid method adopted by Schmedes and Homer (1989). The TBARS value was expressed as milligram malonaldehyde per kilogram of mutton sample.

### Microbial assessment

Microbial assessments such as total viable count, coliform and yeast-mould count were determined. Approximately 10 g of mutton sample was aseptically exercised from stored stock sample. Each stored meat samples was aseptically exercised. Each stored meat sample was thoroughly and uniformly blended in a mechanical blender using sterile diluents (0.1% peptone water) as per recommendation of International Organization for Standardization. About 10g of ground mutton sample was taken aseptically and transferred to a sterile container that contains 90 ml of 0.1% peptone water. Then a homogenized suspension was prepared with a sterile blender and thus 1:10 dilution of sample was produced. A whirly mixture machine of different serial dilutions ranging from  $10^{-2}$  to  $10^{-6}$  were counted through a colony counter. Average number of colonies in a particular dilution was multiplied by the dilution factor to obtain total coliform count. Total coliform count was calculated according to ISO (1995). The results of total coliform count were expressed as the number of an organism of colony forming units per gram (CFU/g) of mutton sample.

### Statistical Analysis

The used model for the planned experiment was factorial experiment with two factors A (Treatments) and B (Days of Intervals) is:

$$Y_{ijk} = \mu + A_i + B_j + (AB)_{ij} + e_{ijk} \quad i=1, \dots, a; j=1, \dots, b; k=1, \dots, n$$

Where:

$Y_{ijk}$  = observation k in level i of factor A and level j of factor B

$\mu$  = the overall mean

$A_i$  = the effect of level i of factor A

$B_j$  = the effect of level j of factor B

Data were analyzed using SAS Statistical Discovery software, NC, USA. Duncan's Multiple Range Test (DMRT) was used to assess the differences among treatments.

## Results and Discussion

### Sensory Evaluation

#### Color

The values of color of the sample are presented in the Table 1. Gamma Irradiation increased the color of mutton significantly ( $P < 0.05$ ). The range of overall observed color score at different doses of irradiation was 3.33 to 4.33. The color value was decreased significantly with storage time and the range was 4.66 to 3.25 ( $P < 0.05$ ). Kim et al. (2012) stated that the redness of meat increased with the formation of carbon monoxide-myoglobin (CO-Mb) complexes. The CO-Mb complex is not easily oxidized to brown metmyoglobin compared with oxymyoglobin due to the strong binding of CO to the iron-porphyrin in myoglobin molecule. They also stated that gamma-ray increased the redness of pork sausages at higher at higher dose level (Ham et al., 2017). Similar results also found by Miller et al. (1995) that irradiation increased the intensity of the color of meat. Nam et al. (2003) observed that different mechanisms and pigments were responsible for irradiation-induced color changes in meat and meat products.

#### Flavor

The score of mean flavors of irradiated and non-irradiated with treatments and days interval are shown in Table 1. The flavor was significantly decreased with treatment (range of flavor score was 4.33 to 3.33) ( $P < 0.05$ ). The flavor was significantly ( $P < 0.05$ ) decreased with days interval and the range of flavor among different days interval was 4.33 to 3.12. Similar results were also found by Modi et al. (2008) where they revealed changes in a goat meat flavor due to gamma irradiation. Irradiation is also known to produce off-odors and off-flavors in food products including beef by the formation of radiolysis products that are dependent on radiation dose, dose rate, packaging conditions and temperature (Jung, 2007).

#### Tenderness

Tenderness in both irradiated and non-irradiated mutton with different days of intervals is shown in Table 1. The range of values of tenderness score at different treatments was 3.66 to 4.11 and there was no significant effect of irradiation on tenderness of mutton ( $P > 0.05$ ). Tenderness was significantly decreased with storage time (range was 4.58 to 3.25) ( $P < 0.05$ ). Similar results were also found in beef in the different study of the same research group Haque et al. (2017). Sweetie et al. (2015) reported that collagen is an abundant tissue protein that plays a significant role in the tenderness and texture of meat. Cheung et al. (1990) found a significant damage of collagen fibers at a higher dose of gamma irradiation. Moreover, molecular weight was also significantly reduced due to cleavage of peptide bonds through radiation which could cause considerable changes in the long-term characteristics of meat. The cross linkage of collagen, actin and reticulin in the endomycium, perimycium and epimycium are responsible for the toughness of meat. Therefore, gamma irradiation might be a suitable tool to prepare tenderer meat through break down of collagen cross linkage.

Table 1. Effect of irradiation on sensory attributes of mutton (mean  $\pm$  SE)

Parameter	DI	Treatment				Mean $\pm$ SE	Level of significance		
		T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	T <sub>4</sub>		T	DI	T $\times$ DI
Colour	0	4.33 $\pm$ 0.33	4.66 $\pm$ 0.33	4.66 $\pm$ 0.33	5.00 $\pm$ 0.00	4.66 <sup>a</sup> $\pm$ 0.24	0.0020	<.0001	0.7000
	30	3.33 $\pm$ 0.33	4.33 $\pm$ 0.33	4.66 $\pm$ 0.33	4.33 $\pm$ 0.33	4.16 <sup>b</sup> $\pm$ 0.33			
	60	2.33 $\pm$ 0.33	3.33 $\pm$ 0.33	3.66 $\pm$ 0.33	3.66 $\pm$ 0.33	3.25 <sup>c</sup> $\pm$ 0.33			
	Mean	3.33 <sup>b</sup> $\pm$ 0.33	4.11 <sup>a</sup> $\pm$ 0.33	4.33 <sup>a</sup> $\pm$ 0.33	4.33 <sup>a</sup> $\pm$ 0.22				
Flavour	0	4.66 $\pm$ 0.33	4.33 $\pm$ 0.33	4.33 $\pm$ 0.33	4.00 $\pm$ 0.57	4.33 <sup>a</sup> $\pm$ 0.39	0.0116	0.0002	0.6537
	30	4.33 $\pm$ 0.33	4.00 $\pm$ 0.00	3.33 $\pm$ 0.33	3.66 $\pm$ 0.33	3.83 <sup>a</sup> $\pm$ 0.24			
	60	4.00 $\pm$ 0.00	3.16 $\pm$ 0.16	3.00 $\pm$ 0.57	2.33 $\pm$ 0.33	3.12 <sup>b</sup> $\pm$ 0.26			
	Mean	4.33 <sup>a</sup> $\pm$ 0.22	3.83 <sup>ab</sup> $\pm$ 0.16	3.55 <sup>b</sup> $\pm$ 0.41	3.33 <sup>b</sup> $\pm$ 0.41				
Tenderness	0	4.33 $\pm$ 0.33	5.00 $\pm$ 0.00	4.66 $\pm$ 0.33	4.33 $\pm$ 0.66	4.58 <sup>a</sup> $\pm$ 0.33	0.6810	0.0018	0.8281
	30	3.66 $\pm$ 0.33	4.00 $\pm$ 0.00	4.33 $\pm$ 0.33	3.66 $\pm$ 0.66	3.91 <sup>ab</sup> $\pm$ 0.33			
	60	3.00 $\pm$ 0.57	3.33 $\pm$ 0.33	3.00 $\pm$ 0.57	3.66 $\pm$ 0.66	3.25 <sup>b</sup> $\pm$ 0.53			
	Mean	3.66 <sup>a</sup> $\pm$ 0.41	4.11 <sup>a</sup> $\pm$ 0.11	4.00 <sup>a</sup> $\pm$ 0.41	3.88 <sup>a</sup> $\pm$ 0.66				
Juiciness	0	4.00 $\pm$ 0.00	4.66 $\pm$ 0.33	4.66 $\pm$ 0.33	5.00 $\pm$ 0.00	4.58 <sup>a</sup> $\pm$ 0.16	0.2007	<.0001	0.9004
	30	3.66 $\pm$ 0.66	3.66 $\pm$ 0.33	4.33 $\pm$ 0.33	4.33 $\pm$ 0.33	4.00 <sup>b</sup> $\pm$ 0.41			
	60	3.00 $\pm$ 0.57	3.00 $\pm$ 0.57	3.00 $\pm$ 0.00	3.33 $\pm$ 0.33	3.08 <sup>c</sup> $\pm$ 0.36			
	Mean	3.55 <sup>a</sup> $\pm$ 0.41	3.77 <sup>a</sup> $\pm$ 0.41	4.00 <sup>a</sup> $\pm$ 0.22	4.22 <sup>a</sup> $\pm$ 0.22				
Overall Acceptability	0	4.33 $\pm$ 0.33	4.66 $\pm$ 0.33	4.66 $\pm$ 0.33	4.66 $\pm$ 0.33	4.58 <sup>a</sup> $\pm$ 0.33	0.6843	0.0027	0.8960
	30	3.66 $\pm$ 0.33	3.33 $\pm$ 0.33	4.00 $\pm$ 0.00	4.16 $\pm$ 0.44	3.79 <sup>b</sup> $\pm$ 0.27			
	60	3.33 $\pm$ 0.88	3.66 $\pm$ 0.33	3.33 $\pm$ 0.33	3.66 $\pm$ 0.33	3.50 <sup>b</sup> $\pm$ 0.46			
	Mean	3.77 <sup>a</sup> $\pm$ 0.51	3.88 <sup>a</sup> $\pm$ 0.33	4.00 <sup>a</sup> $\pm$ 0.22	4.16 <sup>a</sup> $\pm$ 0.36				

Means in each row having different superscripts vary significantly at values  $P < 0.05$ . Again, mean values having same superscript in each row did not differ significantly at  $P > 0.05$ . T<sub>1</sub>=Controlled group, T<sub>2</sub>= 1.5kGy irradiated group, T<sub>3</sub>= 2 kGy irradiated group T<sub>4</sub>= 4 kGy irradiated group, DI=Days of Interval, T= Treatment, T $\times$ DI=Interaction of Treatment and Days of Interval.

Table 2. Effect of irradiation of proximate component of mutton (mean  $\pm$  SE)

Parameter	DI	Treatment				Mean $\pm$ SE	Level of significance		
		T <sub>0</sub>	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>		T	DI	T $\times$ DI
DM (%)	0	20.92 $\pm$ 0.008	21.83 $\pm$ 0.009	22.66 $\pm$ 0.009	23.53 $\pm$ 0.011	22.23 <sup>c</sup> $\pm$ 0.010	<.0001	<.0001	<.0001
	30	21.91 $\pm$ 0.020	22.81 $\pm$ 0.003	23.76 $\pm$ 0.009	24.56 $\pm$ 0.011	23.26 <sup>b</sup> $\pm$ 0.010			
	60	22.28 $\pm$ 0.005	23.25 $\pm$ 0.005	24.08 $\pm$ 0.006	24.91 $\pm$ 0.015	23.63 <sup>a</sup> $\pm$ 0.008			
	Mean	21.70 <sup>d</sup> $\pm$ 0.011	22.63 <sup>c</sup> $\pm$ 0.006	23.50 <sup>b</sup> $\pm$ 0.008	24.33 <sup>a</sup> $\pm$ 0.012				
CP (%)	0	21.70 $\pm$ 0.005	21.73 $\pm$ 0.005	21.69 $\pm$ 0.011	21.61 $\pm$ 0.006	21.68 <sup>a</sup> $\pm$ 0.006	0.0057	<.0001	<.0001
	30	19.64 $\pm$ 0.012	19.67 $\pm$ 0.017	19.60 $\pm$ 0.012	19.62 $\pm$ 0.015	19.72 <sup>b</sup> $\pm$ 0.014			
	60	17.43 $\pm$ 0.011	17.46 $\pm$ 0.005	17.48 $\pm$ 0.009	17.48 $\pm$ 0.012	17.46 <sup>c</sup> $\pm$ 0.009			
	Mean	19.59 <sup>ab</sup> $\pm$ 0.009	19.61 <sup>a</sup> $\pm$ 0.009	19.58 <sup>b</sup> $\pm$ 0.010	19.57 <sup>b</sup> $\pm$ 0.011				
EE (%)	0	12.50 $\pm$ 0.020	13.13 $\pm$ 0.011	13.64 $\pm$ 0.017	14.22 $\pm$ 0.032	13.37 <sup>c</sup> $\pm$ 0.02	<.0001	<.0001	<.0001
	30	12.94 $\pm$ 0.017	13.83 $\pm$ 0.008	14.74 $\pm$ 0.020	15.60 $\pm$ 0.026	14.28 <sup>b</sup> $\pm$ 0.017			
	60	14.26 $\pm$ 0.031	15.10 $\pm$ 0.020	15.97 $\pm$ 0.008	16.85 $\pm$ 0.024	15.54 <sup>a</sup> $\pm$ 0.020			
	Mean	13.23 <sup>d</sup> $\pm$ 0.022	14.02 <sup>c</sup> $\pm$ 0.013	14.78 <sup>b</sup> $\pm$ 0.015	15.56 <sup>a</sup> $\pm$ 0.027				
ASH (%)	0	2.02 $\pm$ 0.093	1.94 $\pm$ 0.075	1.82 $\pm$ 0.053	1.70 $\pm$ 0.049	1.87 <sup>a</sup> $\pm$ 0.067	<.0001	0.0453	0.9967
	30	2.00 $\pm$ 0.093	1.88 $\pm$ 0.091	1.75 $\pm$ 0.085	1.62 $\pm$ 0.070	1.81 <sup>ab</sup> $\pm$ 0.084			
	60	1.95 $\pm$ 0.028	1.81 $\pm$ 0.030	1.67 $\pm$ 0.037	1.56 $\pm$ 0.033	1.75 <sup>b</sup> $\pm$ 0.032			
	Mean	1.99 <sup>a</sup> $\pm$ 0.071	1.88 <sup>b</sup> $\pm$ 0.065	1.75 <sup>c</sup> $\pm$ 0.058	1.63 <sup>d</sup> $\pm$ 0.050				

Means in each row having different superscripts vary significantly at values  $P < 0.05$ . Again, mean values having same superscript in each row did not differ significantly at  $P > 0.05$ . T<sub>0</sub>=Control group, T<sub>1</sub>= 1.5 kGy irradiated group, T<sub>2</sub>= 2 kGy irradiated group T<sub>3</sub>= 4 kGy irradiated group, DI=Days of Interval, T= Treatment, T $\times$ DI=Interaction of Treatment and Days of Interval.

### Juiciness

Mean scores in both irradiated and non-irradiated mutton have shown in Table 1. The values of juiciness at different treatments were ranges 3.55 to 4.22 and there was no significant effect of irradiation on the juiciness values of mutton ( $P > 0.05$ ). Juiciness was significantly decreased and the range of juiciness score was 4.58 to 3.08 with storage time ( $P < 0.05$ ). Similarly, Badr (2004) found that irradiation in rabbit meat had no significant effect on the juiciness. Miyagasku et al. (2003) examined chicken breast meat and stated that 3.0 kGy level of irradiation dosage increases the shelf life from 5 to 22 days without modifying the sensory parameters.

### Overall Acceptability

The overall acceptability values at different treatments were ranges 3.77 to 4.16. Irradiation doses slightly increased overall acceptability of meat but it was not significant ( $P > 0.05$ ). As well, Johnson et al. (2004) found significant difference between the irradiated and non-irradiated meat. The range of overall acceptability score was 4.58 to 3.50 with different days of interval. The overall acceptability significantly decreased with storage time ( $P < 0.05$ ). Johnson et al. (2004) also found that overall acceptance, acceptance of flavor, juiciness, tenderness and mouth feeling of the non-irradiated chicken and its products were significantly lower than irradiated samples

at day 18 and day 32 after irradiation, respectively ( $P<0.05$ ). Haque et al. (2017) found the similar results of overall acceptability in beef.

### **Proximate Analysis**

Effects of gamma irradiation on the proximate composition of mutton are presented in Table 2. Apparently, irradiation treatments brought only minor changes in the DM, CP, EE and ash contents of meats.

#### **Dry Matter (DM)**

DM content of mutton was increased significantly with irradiation ( $P<0.05$ ). The range of DM content at different treatments was 21.70% to 24.33%. DM content also increased and the range was 22.23% to 23.63% with storage time ( $P<0.05$ ). Extra cellular fluid of meat might be shrunk during irradiation process, resulting in less moisture content in the muscle. As the irradiation breaks the collagen and other fibers might help to reduce or evaporate water from muscle after irradiation (Cheung et al., 1990). Al-Bachir and Zeinou (2009) and Modi et al. (2008) were also found the similar patterns in which moisture percentage was decreased with increasing irradiation dosage and storage time. A similar pattern was also observed by Konieczny et al. (2007) where they reported that DM content increased with the advancement of storage time.

#### **Crude Protein (CP)**

The range of CP% at different treatment level was 19.57 to 19.61 %. Results showed that the highest level ( $P<0.05$ ) of CP% was found in 1.5 kGy irradiated group than that of other treatment group. Unlikely, Modi et al. (2008), Al-Bachir and Zeinou (2009) and Haque et al. (2017) found that CP content of meat did not significantly change due to irradiation. The CP content significantly decreased with storage time which ranges from 21.68 to 17.46% ( $p<0.05$ ) and these results are in agreement with Al-Bachir and Zeinou (2009) where CP content decreased with increasing storage time in meat and meat products.

#### **Ether Extract (EE)**

Irradiated samples showed significantly higher amounts of EE compared with controlled group ( $P<0.05$ ). The effect of the gamma irradiation on the fatty acid composition can be ascribed to the production of free radicals during the irradiation of mutton. Water losses occurred during irradiation that resulting a higher DM content in irradiated mutton than in non-irradiated might be the cause of higher lipid content in most cases. This result may be explained by the inverse relationship between fat and moisture contents or a dose-dependent effect of irradiation on the fatty acid profile (Stefanova et al., 2011). The higher amount fat content might be due to the degradation of large lipid molecules which ultimately added to the fat content of the sample. This trend was similar to that of (Al-Bachir and Zeinou, 2009; Yilmaz and Gecgel, 2007; Fernandez-Lopez et al., 2003). The EE content (13.37 to 15.54%) also significantly increased with storage time ( $P<0.05$ ). The results in this study were very close with the findings of Al-Bachir and Zeinou (2009) and Bakalivanova et al. (2009).

#### **Ash**

The ash content of both irradiated and non-irradiated meat was shown in Table 2. Ash content was significantly decreased with the increasing of irradiation dosage

( $P<0.05$ ) but insignificant in days of intervals as well as interaction of treatment and days of intervals. Decreasing of ash content after irradiation might be due to the physical disruption of muscle fibers and also increasing of oxidation and peroxide values during irradiation process. Al-Bachir and Zeinou (2009) stated that ash content of meat slightly increased with different irradiation dosage as well as days of intervals but these were not significant. Gecgel (2013) reported similar findings for meatballs, but their samples were irradiated at doses up to 7.0 kGy. Arannilewa et al. (2005) observed that ash content of the meat is decreased with frozen storage. A similar result was also found by Haque et al. (2017) in beef.

### **Physicochemical and Bio-chemical Properties**

The physicochemical and Bio-chemical properties such as raw pH, cooking loss, PV, TBARS, FFA were determined and the results are presented in Table 3.

#### **Raw pH**

The range of raw pH was 5.86% to 5.74% with irradiation dose. The pH value was decreased significantly with increasing doses of irradiation ( $P<0.05$ ). This is because an increase in fat values in irradiated samples caused decrease raw pH of meat (Morales et al. 2009). Raw pH also significantly decreased with storage time due to increased free fatty acids because of rancidity ( $P<0.05$ ). Similar result also found in irradiated broiler meat where the pH was slightly decreased with the increasing of irradiation as well as with the advancement of storage time in refrigerator (Morales et al. 2009). Similarly, Aftab et al. (2015) also found that raw pH was decreased with higher irradiation as well as storage.

#### **Cooking Loss**

The range of cooking loss at different treatments was 38.58% to 40.76%. The results showed that cooking loss was significantly increased with irradiation doses as well as days of interval ( $P<0.05$ ). Irradiation as well as storage time decreased muscle fiber that was the cause of increased cooking losses. Yoon (2003) reported that irradiated chicken breast showed a higher cooking loss than non-irradiated meat also supported the present study.

#### **Peroxide Value (PV-meq/kg)**

The range of overall observed PV at different treatment levels was 0.44 to 0.95. PV values were increased significantly with increasing irradiation doses ( $P<0.05$ ). The PV values also increased significantly with storage time and the overall observed values were 0.54 to 0.65 ( $p<0.05$ ). Similarly, the PV was increased in beef with the increased irradiation dosage (Haque et al., 2017; Quattara et al., 2002).

#### **Thiobarbituric Acid Value (TBARS)**

The range of TBARS value of different treatment was 0.42 to 0.61. The TBARS values were increased significantly with increased treatment doses and storage time ( $P<0.05$ ). The results of the present study are very much similar with the few other researches (Al-Bachir and Zeinou, 2009; Bakalivanova et al., 2009). They reported an increase in oxidation activity and lipid peroxidation as a result of both irradiation treatment and storage time on meat and meat products. Kanatt et al. (2006) showed that TBA value increased during irradiation and storage in lamb meat.

Table 3. Effect of irradiation on physicochemical and bio-chemical properties of mutton (mean  $\pm$  SE)

Parameter	DI	Treatment				Mean $\pm$ SE	Level of significance		
		T <sub>0</sub>	T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>		T	DI	T $\times$ DI
Raw pH	0	5.91 $\pm$ 0.005	5.87 $\pm$ 0.005	5.82 $\pm$ 0.014	5.78 $\pm$ 0.017	5.84 <sup>a</sup> $\pm$ 0.010			
	30	5.85 $\pm$ 0.006	5.81 $\pm$ 0.014	5.77 $\pm$ 0.017	5.74 $\pm$ 0.011	5.79 <sup>b</sup> $\pm$ 0.012	<.0001	<.0001	0.9635
	60	5.82 $\pm$ 0.005	5.80 $\pm$ 0.005	5.74 $\pm$ 0.011	5.71 $\pm$ 0.020	5.76 <sup>c</sup> $\pm$ 0.010			
	Mean	5.86 <sup>a</sup> $\pm$ 0.005	5.82 <sup>b</sup> $\pm$ 0.008	5.77 <sup>c</sup> $\pm$ 0.014	5.74 <sup>d</sup> $\pm$ 0.016				
Cooking Loss	0	34.66 $\pm$ 0.018	35.45 $\pm$ 0.020	36.16 $\pm$ 0.008	37.03 $\pm$ 0.011	35.82 <sup>c</sup> $\pm$ 0.014			
	30	38.94 $\pm$ 0.020	39.73 $\pm$ 0.008	40.36 $\pm$ 0.008	41.12 $\pm$ 0.012	40.04 <sup>b</sup> $\pm$ 0.012	<.0001	<.0001	0.2012
	60	42.15 $\pm$ 0.017	42.86 $\pm$ 0.008	43.47 $\pm$ 0.008	44.15 $\pm$ 0.023	43.16 <sup>a</sup> $\pm$ 0.014			
	Mean	38.58 <sup>d</sup> $\pm$ 0.018	39.35 <sup>c</sup> $\pm$ 0.012	40.00 <sup>b</sup> $\pm$ 0.008	40.76 <sup>a</sup> $\pm$ 0.015				
PV	0	0.35 $\pm$ 0.039	0.43 $\pm$ 0.008	0.52 $\pm$ 0.015	0.88 $\pm$ 0.012	0.54 <sup>b</sup> $\pm$ 0.018			
	30	0.48 $\pm$ 0.023	0.53 $\pm$ 0.017	0.57 $\pm$ 0.020	0.95 $\pm$ 0.017	0.63 <sup>a</sup> $\pm$ 0.019	<.0001	<.0001	0.0935
	60	0.48 $\pm$ 0.012	0.55 $\pm$ 0.011	0.57 $\pm$ 0.011	1.03 $\pm$ 0.011	0.65 <sup>a</sup> $\pm$ 0.011			
	Mean	0.44 <sup>d</sup> $\pm$ 0.024	0.50 <sup>c</sup> $\pm$ 0.012	0.55 <sup>b</sup> $\pm$ 0.015	0.95 <sup>a</sup> $\pm$ 0.013				
TBARS	0	0.14 $\pm$ 0.011	0.19 $\pm$ 0.011	0.26 $\pm$ 0.011	0.35 $\pm$ 0.014	0.23 <sup>c</sup> $\pm$ 0.011			
	30	0.45 $\pm$ 0.014	0.51 $\pm$ 0.014	0.56 $\pm$ 0.014	0.65 $\pm$ 0.014	0.54 <sup>b</sup> $\pm$ 0.014	<.0001	<.0001	0.4708
	60	0.68 $\pm$ 0.017	0.71 $\pm$ 0.017	0.75 $\pm$ 0.014	0.83 $\pm$ 0.014	0.74 <sup>a</sup> $\pm$ 0.015			
	Mean	0.42 <sup>d</sup> $\pm$ 0.014	0.47 <sup>c</sup> $\pm$ 0.014	0.52 <sup>b</sup> $\pm$ 0.013	0.61 <sup>a</sup> $\pm$ 0.014				
FFA	0	0.61 $\pm$ 0.017	0.78 $\pm$ 0.023	0.87 $\pm$ 0.008	0.91 $\pm$ 0.020	0.79 <sup>c</sup> $\pm$ 0.017			
	30	0.71 $\pm$ 0.020	0.85 $\pm$ 0.011	0.97 $\pm$ 0.032	1.08 $\pm$ 0.017	0.90 <sup>b</sup> $\pm$ 0.02	<.0001	<.0001	<.0001
	60	0.89 $\pm$ 0.017	1.06 $\pm$ 0.029	1.12 $\pm$ 0.024	1.25 $\pm$ 0.020	1.08 <sup>a</sup> $\pm$ 0.022			
	Mean	0.74 <sup>d</sup> $\pm$ 0.018	0.90 <sup>c</sup> $\pm$ 0.021	0.99 <sup>b</sup> $\pm$ 0.015	1.08 <sup>a</sup> $\pm$ 0.019				

Means in each row having different superscripts vary significantly at values  $P < 0.05$ . Again, mean values having same superscript in each row did not differ significantly at  $P > 0.05$ . T<sub>1</sub>=Controlled group, T<sub>2</sub>= 1.5 kGy irradiated group, T<sub>3</sub>= 2 kGy irradiated group T<sub>4</sub>= 4 kGy irradiated group, DI=Days of Interval, T= Treatment, T $\times$ DI=Interaction of Treatment and Days of Interval.

Table 4. Effect of different doses of irradiation on microbial quality (mean  $\pm$  SE) of mutton

Parameter	DI	Treatment				Mean $\pm$ SE	Level of significance		
		T <sub>1</sub>	T <sub>2</sub>	T <sub>3</sub>	T <sub>4</sub>		T	DI	T $\times$ DI
TVC (logCFU/g)	0	5.61 $\pm$ 0.032	5.06 $\pm$ 0.050	4.55 $\pm$ 0.063	3.65 $\pm$ 0.068	4.72 <sup>c</sup> $\pm$ 0.053			
	30	5.89 $\pm$ 0.027	5.13 $\pm$ 0.043	4.67 $\pm$ 0.139	3.76 $\pm$ 0.150	4.86 <sup>b</sup> $\pm$ 0.065	<.0001	<.0001	0.1133
	60	6.38 $\pm$ 0.018	5.30 $\pm$ 0.125	4.95 $\pm$ 0.054	4.06 $\pm$ 0.050	5.17 <sup>a</sup> $\pm$ 0.061			
	Mean	5.96 <sup>a</sup> $\pm$ 0.025	5.16 <sup>b</sup> $\pm$ 0.072	4.73 <sup>c</sup> $\pm$ 0.043	3.82 <sup>d</sup> $\pm$ 0.089				
TCC (logCFU/g)	0	2.03 $\pm$ 0.038	1.34 $\pm$ 0.024	1.14 $\pm$ 0.050	0.95 $\pm$ 0.021	1.36 <sup>c</sup> $\pm$ 0.033			
	30	2.57 $\pm$ 0.056	2.03 $\pm$ 0.069	1.75 $\pm$ 0.026	1.37 $\pm$ 0.043	1.93 <sup>b</sup> $\pm$ 0.048	<.0001	<.0001	0.0031
	60	2.92 $\pm$ 0.015	2.33 $\pm$ 0.074	1.83 $\pm$ 0.029	1.55 $\pm$ 0.055	2.16 <sup>a</sup> $\pm$ 0.043			
	Mean	2.51 <sup>a</sup> $\pm$ 0.036	1.90 <sup>b</sup> $\pm$ 0.055	1.57 <sup>c</sup> $\pm$ 0.035	1.29 <sup>d</sup> $\pm$ 0.039				
TYMC (logCFU/g)	0	3.13 $\pm$ 0.009	2.56 $\pm$ 0.054	2.18 $\pm$ 0.043	1.87 $\pm$ 0.050	2.43 <sup>c</sup> $\pm$ 0.039			
	30	3.61 $\pm$ 0.180	2.78 $\pm$ 0.033	2.42 $\pm$ 0.046	1.92 $\pm$ 0.040	2.68 <sup>b</sup> $\pm$ 0.074	<.0001	<.0001	0.0002
	60	4.41 $\pm$ 0.003	3.15 $\pm$ 0.058	2.75 $\pm$ 0.060	2.47 $\pm$ 0.058	3.20 <sup>a</sup> $\pm$ 0.044			
	Mean	3.72 <sup>a</sup> $\pm$ 0.064	2.83 <sup>b</sup> $\pm$ 0.048	2.45 <sup>c</sup> $\pm$ 0.049	2.09 <sup>d</sup> $\pm$ 0.049				

Means in each row having different superscripts vary significantly at values  $p < 0.05$ . Again, mean values having same superscript in each row did not differ significantly at  $p > 0.05$ . T<sub>1</sub>=Controlled group, T<sub>2</sub>= 1.5 kGy irradiated group, T<sub>3</sub>=2 kGy irradiated group, T<sub>4</sub>= 4 kGy irradiated group, DI=Days of Interval, T= Treatment, T $\times$ DI=Interaction of Treatment and Days of Interval.

#### Free Fatty Acid Value (FFA %)

The range of overall observed FFA of different treatments was 0.74 to 1.08%. The FFA results appeared to be consistent with those of thiobarbituric acid reactive substances (TBA) and peroxide value (POV). The FFA values are significantly increased with increased irradiation dose and storage time ( $P < 0.05$ ). Rima et al. (2019) stated that irradiated chicken with 4.5 kGy dose level showed a higher FFA content compared to non-irradiated control samples. Similarly, Chen et al. (2007) found that percentage of saturated fatty acids increased significantly with storage time and after the irradiation.

#### Microbiological Assessment

In this study, the effect of gamma irradiation on microflora (TVC) and food borne pathogens (Coliform and Yeast-Mold) on both controlled and irradiated mutton were observed. The results are presented in Table 4.

#### Total viable count (TVC)

The range observed of TVC with different treatments value was 5.96 to 3.82 (log CFU/gm). The TVC value was decreased significantly with irradiation dose ( $P < 0.05$ ). The range of TVC value was 4.72 to 5.17 with different days of intervals. The TVC value was significantly increased with storage time ( $P < 0.05$ ). Ferawati et al. (2015) reported a lower total plate count in the irradiated samples compared to control. Therefore, it could be said that irradiation is a way of reducing microbial load that might be helpful to increase shelf life of fresh mutton.

#### Total coliform count (TCC)

Results showed that TCC value was decreased significantly with irradiation dose ( $P < 0.05$ ). The range of observed TCC with different treatments was 2.51 to 1.29 (log CFU/gm). The TCC values were significantly increased with storage time ( $P < 0.05$ ). Inamura et al. (2012) reported that irradiated samples showed decrease in TCC and might be safe up to 8 months of storage after gamma irradiation.

### Total yeast-molds count (TYMC)

The TYMC value was decreased significantly with treatment (3.27 to 2.09 log CFU/gm) ( $p < 0.05$ ), but significantly increased with storage time (2.43 to 3.20 log CFU/gm) ( $p < 0.05$ ). The lower value of TYMC indicates that this product is suitable for consumer's health. Similarly, Haque et al. (2017) showed that irradiation significantly decreased yeast and mold count of beef ( $p < 0.05$ ). Ahmed et al. (2009) stated that 4 kGy irradiation levels were suitable for controlling fungal growth on sun dried fish. Badr (2004) reported that the yeasts and molds significantly reduced (84 and 94%, respectively) by the irradiation of rabbit meat. Haque et al. (2017) stated that 6 kGy irradiation dose showed the most acceptable for microbial population reduction, maintains overall acceptability and longer shelf life during beef preservation compared with 2 and 4 kGy irradiation dosages.

### Conclusions

The results of the present study clearly demonstrated that gamma irradiation increased color, tenderness, juiciness and overall acceptability of mutton. Different doses of irradiation increased the shelf life of meat without changing nutritional and sensory properties of meat. Among the treatments, 4.0 kGy showed the best results in terms of sensory attributes, redox control, and microbial population reduction. The main objective of irradiation of meat is to increase the shelflife and to remove the international trade barrier where disease is factors. More study should be needed on specific microorganisms related to spoilage and safety of meats.

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### Ethics Approval

This article does not require IRB/IACUC approval because there are no human and animal participants

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