

## EFFECT OF GAMMA IRRADIATION ON SOME MICROORGANISMS CONTAMINATING ANIMAL FEEDS

R.M. BADAWAY, M.M. ZENHOM

And A.I. HAMMAD

Food Technology Research Institute , Agricultural Research Centre, Giza, Egypt.

(Manuscript received 7 December 1993)

### Abstract

Microbial determination in animal feeds produced in Egypt or fish meal imported from abroad was carried out Gamma irradiation with the dose 20KGY sterilized the pathogens, while 5 KGY destroyed them all. The microbes experimented were sporeformers, yeast and molds, *Staphylococcus aureus* and *Streptococcus faecalis*.

### INTRODUCTION

The need for animal feeds has increased with the expansion of animal farms and the deficiency in animal protein production in Egypt. It has been reported however that animal feeds and their components especially fish, meat-bone and blood meals are the main factors of contamination with pathogenic microorganisms. These pathogens can cause diseases in poultry and other animals (Smith, 1970). For this reason great attention must be given to high quality animal feed.

Sterilisation of this feed can be achieved by heating which might also cause destruction of feed components (vitamines and/or proteins). Also, it is not always easy or feasible to destroy pathogens in horn meal and crushed bones (Egyum, 1969). Fumigation by chemicals also causes health hazard to workers and toxins still remained. Gamma irradiation was found to require low energy in comparison with heating (Hansen, 1966; Ley *et al.*, 1969).

---

(\*) : National Centre for Radiation Research & Technology, Atomic Energy Authority, Cairo-Egypt.

Farag (1982) found that the initial microbial counts were  $4.4 \times 10^5$ ,  $2.1 \times 10^4$  and  $7.0 \times 10^3$  cells / gm for blood, meat and fish meals, respectively. El-Zawahry *et al.* (1985) found that total bacterial counts ranged between  $2.2 \times 10^3$  and  $2.1 \times 10^4$  cells / gm for 3 examined poultry feed samples. The isolate species belonged to families *Bacillaceae*, *Micrococcaceae*, *Pseudomonadaceae* and *Enterobacteriaceae*.

Farage, (1982) found that irradiation dose of 5.0 KGY reduced the initial bacterial counts of blood meal by 2 log cycles, by one log cycle for the counts of meat, and destroyed all the counts of fish meal. El-Zawahry *et al.* (1985) reported that 20 KGY were required for complete inhibition of poultry feed flora.

## MATERIALS AND METHODS

### Sampling

Soybean meal was secured from the Egyptian Company for Poultry. Fish meal was imported. Meat-bone meal was secured from the Mechanical Slaughter, Gerco Company (El-Basatein). Blood meal was obtained from the organic Fertilizers Company. Three samples from the aforementioned types were left for comparison. The samples were exposed to the dosages 5.0, 10.0, 12.5 and 20.0 KGY of gamma irradiation, and the dose rate of the source was 0.08 KGY/SEC. The irradiation facility used was the Egyptian Industrial Mega Gamma Irradiation.

### Microbiological estimations

30 g of ground sample were homogenized in 270 ml of 0.1% peptone water for 5 minutes in a sterile blender jar. Serial dilutions up to  $10^8$  of the homogenate were used. Aerobic plate count and aerobic sporeformers after pasteurisation at 80°C for 15 minutes were done on standard plate count agar and incubated at 37°C for 48 h. Yeast and mold were counted on malt extract agar (Difco Manual, 1977) at 25°C for 5 days. *Staphylococcus aureus* was grown on staphylococcus medium No. 110 (Difco, 1977) and *Strep. faecalis* on Kanamycin aesculine azide agar (Oxoid Manual, 1982). Both species were grown at 37°C for 24 h.

## RESULTS AND DISCUSSION

Data tabulated in Table 1 show that the dose of 5.0 KGY gamma irradiation had reduced the initial total bacterial counts between one and three log cycles. The

highest reduction occurred in meat-bone meal, while intermediate reduction occurred in blood and fish meals. Counts in soybean however, had shown the highest resistance to this dose. Increasing the dose to 10.0 KGY increased the reduction by nearly 1-2 log cycle, while the dose 12.5 increased the reduction by two log cycles. The dose 20.0 KGY completely reduced microbes detected in the initial counts. This is in agreement with the findings of Aziz (1982) and El-Zawahry *et al.* (1985) who reported a reduction by 1-210g cycle for 5.0 KGY and a severe reduction with the doses ranging between 10.0-20.0 KGY.

Table 1 indicates that the aerobic sporeforming bacteria was the most resistant species among the other examined microorganisms. This is in accordance with the results of Ito *et al.* (1985). The reduction by different doses almost showed the same trend observed with the total bacterial counts, since the application of 12.5 KGY nearly had a lethal effect with aerobic sporeformers.

As shown in Table 1, yeast and mold were completely destroyed with the doses 12.5 and 20.0 KGY. On the other hand 5.0 KGY was sufficient to destroy the examined pathogens. This is in agreement with the findings of Ley *et al.* (1969), Ito *et al.* (1981).

Table 1. Effect of gamma irradiation on microbial counts of animal feed components (counts/gm)

Animal feed components	irradiation dose (KGY)				
	0.0	55.0	10.0	12.5	20.0
Total microbial counts					
Soybean meal	7.9x10 <sup>6</sup>	3.5x10 <sup>5</sup>	1.5x10 <sup>3</sup>	4.8x10 <sup>2</sup>	0.0
Fish meal	4.5x10 <sup>6</sup>	2.6x10 <sup>3</sup>	2.0x10 <sup>2</sup>	0.0	0.0
Meat-bone meal	7.0x10 <sup>4</sup>	7.0x10 <sup>2</sup>	4.0x10 <sup>2</sup>	0.0	0.0
Blood meal	3.0x10 <sup>5</sup>	3.5x10 <sup>3</sup>	3.0x10 <sup>2</sup>	0.0	0.0
Sporeformers					
Soybean meal	1.3x10 <sup>6</sup>	3.6x10 <sup>3</sup>	6.0x10 <sup>2</sup>	1.5x10 <sup>1</sup>	0.0
Fish meal	2.0x10 <sup>4</sup>	1.0x10 <sup>3</sup>	1.0x10 <sup>2</sup>	0.0	0.0
Meat-bone meal	1.0x10 <sup>3</sup>	2.5x10 <sup>2</sup>	1.5x10 <sup>2</sup>	0.0	0.0
Blood meal	4.0x10 <sup>4</sup>	3.0x10 <sup>3</sup>	6.0x10 <sup>2</sup>	0.0	0.0
Yeast & molds					
Soybean meal	2.2x10 <sup>4</sup>	2.5x10 <sup>2</sup>	4.0x10 <sup>1</sup>	0.0	0.0
Fish meal	3.5x10 <sup>3</sup>	2.0x10 <sup>2</sup>	3.0x10 <sup>1</sup>	0.0	0.0
Meat-bone meal	2.0x10 <sup>3</sup>	2.0x10 <sup>2</sup>	2.0x10 <sup>1</sup>	0.0	0.0
Blood meal	3.0x10 <sup>3</sup>	1.5x10 <sup>2</sup>	2.0x10 <sup>1</sup>	0.0	0.0
<i>Staphylococcus aureus</i>					
Soybean meal	1.0x10 <sup>4</sup>	0.0	0.0	0.0	0.0
Fish meal	2.0x10 <sup>2</sup>	0.0	0.0	0.0	0.0
Meat-bone meal	2.0x10 <sup>2</sup>	0.0	0.0	0.0	0.0
Blood meal	3.0x10 <sup>2</sup>	0.0	0.0	0.0	0.0
<i>Staphylococcus faecalis</i>					
Soybean meal	1.0x10 <sup>5</sup>	0.0	0.0	0.0	0.0
Fish meal	2.0x10 <sup>2</sup>	0.0	0.0	0.0	0.0
Meat-bone meal	2.5x10 <sup>2</sup>	0.0	0.0	0.0	0.0
Blood meal	2.2x10 <sup>2</sup>	0.0	0.0	0.0	0.0

## REFERENCES

- 1 . Aziz, N.H. 1982. The microflora of poultry diets in relation to human and poultry disease and control by gamma radiation. M.Sc. Thesis, Ain Shams Univ., Faculty of Sci., Cairo , Egypt.
- 2 . Difco Manual 1977. Difco manual of dehydrated cultures media and reagent. 9th ed : 32 Difco laboratories incorporated, Detroit Michigan, U.S.A.
- 3 . Egyum, B.O. 1969. Der einfluss der sterilization auf die protein qualitat von futtermischungen.Z. Tierern-Futtermittelk, 25,204 (C.F. Adamiker, D. 1975).
- 4 . El-Zawahry, Y.A., Y.A. Youssef, and R.M. Roushdy 1985. Radiation deactivation of bacterial flora in some Egyptian poultry feed. Food Irradiation Processing "IAEA, Vienna" 1985.
- 5 . Farag, M.D. 1982. Nutritional evaluation of radiation Induced biochemical changes in animal feed by products. M.Sc. Thesis, Al-Azhar Univ., Faculty of Agri., Cairo-Egypt.
- 6 . Hansen, P.I.E. 1966. Radiation treatment of meat products and animal by - products. Food Irradiation "proc. symp. Karlsruhe, 1966). IAEA, Vienna, Pp. 411.
- 7 . Ito, H.T., M. Kum, M. Takenhisa, and H. Izuka 1981. Distribution of microorganisms in animal feed and their disinfection by radiation. Radiat. Phys. Chem., 18 (3-4) 569-574
- 8 . Ley, F.J., J. Bleby, E.C. Marri, and J.C. Paterson, 1969. Sterilization of laboratory animal diets using gamma irradiation. Leb. Anim., (1969), 2,221-254.
- 9 . Oxoid Manual . 1982. The oxoid manual of culture media ingredients and other laboratory services. Oxoid Limited, Hampshire, England.
- 10 . Smith, J.W. and P.B. Hamilton, 1970. Aflatoxicosis in the broiler chicken. Poultry Sci., 94:207-215.

## تأثير اشعة جاما علي الكائنات الحية الدقيقة الملوثة للاغذية الحيوانية

رضا بدوى ، محمود زينهم ، علي جاد

معهد بحوث تكنولوجيا الأغذية ، مركز البحوث الزراعية - الجيزة.

أجريت دراسة تأثير اشعة جاما بجرعات ١٠,٥ ، ١٢,٥ ، ٢٠ كيلو جراي بمعدل ٠,٠٨ كيلو جراي / ث بغرض ازالة الميكروبات الملوثة للاغذية الحيوانية سواء الميكروبات المرضية أو المتجرثمة أو الخمائر أو الفطريات وذلك لتقدير انسب هذه الجرعات. وقد تمت هذه الدراسة بهدف الحصول علي لحوم حيوانات او دواجن خالية من الميكروبات المرضية يسهل استخدامها بعد ذلك في الحصول علي منتجات مصنفة ذات جودة عالية. وكانت عينات مكونات الاعلاف المستخدمة الحبوب مثل مطحون فول الصويا ومخلفات المجازر مثل مسحوق عظام اللحوم ومسحوق الدم ومخلفات تصنيع الأسماك (مسحوق السمك).

وكانت الميكروبات المختبرة هي المتجرثمة الهوائية، والفطريات والخمائر ، والاستافيلوكوكس أوريس ، الاستربتوكوكس فيكالييس ، والعدد الكلي . وأظهرت النتائج ان الجرعة ٢٠ كيلو جراي والتي كانت اقل نسبيا في الفعالية لكنها أزالته تقريبا معظم المجموعة ماعدا الميكروبات المتجرثمة الهوائية ، وأزالته الجرعة ٥٠ كيلو جراي تماما الميكروبات المرضية المختبرة.

وتظهر من الدراسة أن جرعة ٢٠ كيلو جراي لها تأثير معقم علي الميكروبات ويمكن استخدامها في الحصول علي عليقة نظيفة وبالتالي تحقيق لحوم خالية من الميكروبات المرضية.