The effect of ionizing radiation and storage temperature on the post-harvest growth and some quality properties of mushrooms (Agaricus bisporus)

KRYSPINA ŚMIERZCHALSKA and ELŻBIETA WOJNIAKIEWICZ

Research Institute of Vegetable Crops, 22 Lipca 1/3, 96-100 Skierniewice, Poland

(Received: June 14, 1985)

Abstract

The effects of a dose of 2.0 kGy (Co-60) and three ranges of storage temperature (0-4°C, 9-11°C, 18-19°C) on increasing the shelf-life and some quality properties of the mushrooms were investigated. The retardation of mushroom growth and ageing by ionizing radiation was estimated by measurements of the cap diameter, stem elongation, cap opening and discoloration of cap surface. The control of fungal and bacterial diseases was also evaluated. The effect of lower doses, 0.5 and 1.0 kGy, was compared at the storage temperature of 10°C.

The irradiation of mushrooms soon after harvest and storage at temperatures of 10 and 16°C allowed the retention of quality and increased the shelf-life to 8 days; at 18-19°C, to 6 days.

INTRODUCTION

Both increasing production of mushrooms and their perishable properties call for the more economical methods of post-harvest storage.

A review of the experiments carried out in the Netherlands, United States of America, Hungary and Czechoslovakia dealing with the irradiation method has been published (Śmierzchalska and Gubrynowicz, 1979). The results encouraged checking the method under our conditions.

Studies were undertaken in the Isotopic Laboratory at the Department of Biology in 1979. They aimed to show the effectiveness of ionizing radiation on the retardation of mushroom growth and ageing depending on the storage temperature after irradiation. The effect of different doses was also investigated.

In addition to several quality properties, the control of fungal and bacterial diseases by ionizing radiation was also evaluated.

MATERIALS AND METHODS

A number of experiments was carried out during 1979-1981 with mushrooms (*Agaricus bisporus*) to show the effectiveness of ionizing radiation at different storage temperatures. The experimental series included the following treatments:

- irradiation doses of 0 and 2.0 kGy; storage temperatures: $0-4^{\circ}C$, $9-11^{\circ}C$, and $18-19^{\circ}C$,
- irradiation doses of 0 and 2.0 kGy; storage temperatures: 10° C and 16° C,
- irradiation doses of 0, 0.5, 1.0 and 2.0 kGy; storage temperature 10°C.

Each experimental series consisted of 3 or 4 experiments, started at one week intervals. The mushrooms were taken mostly from the first, second and third flushes. In each particular experiment the treatments were done in triplicate.

Mushrooms were delivered from the Experimental and Production Station for Edible Mushrooms in Skierniewice. In the first series, the strain used was Somycel 611, in the next two — Somycel 653. Mushrooms were gathered early in the morning, cleaned, selected and packed into small paper boxes.

Experimental mushrooms were at the "closed cup" stage (unbroken velum) with diameters of 25-40 mm, and stem length of a few millimeters. Twenty-two to thirty mushrooms were put into boxes; their fresh weight amounted to about 200 g. The small boxes were placed into larger boxes in a single layer for transport and storage after irradiation. The small paper boxes, as well as the large ones, were imported from the Netherlands and are used for transportation of mushrooms abroad. When the big boxes with samples were stacked on each other, 90-100% RH was reached inside the small boxes with mushrooms.

Mushroom samples were brought by car to the Institute of Radiation Chemistry in Łódź (70 km from Skierniewice) and irradiated with gamma rays from a Co-60 source with an activity of about 740 TBq (about 20 kCi). To absorb a dose of 2.0 kGy, the irradiation had to last 1 h 40 min. The standard deviations for the applied doses were the following: 2.0 ± 0.2 kGy; 1.0 ± 0.1 kGy; 0.5 ± 0.1 kGy. The temperature in the irradiation chamber was $18-20^{\circ}$ C.

The time between harvest and irradiation was very short; it ranged from 5 to 7 h. During transport the temperature varied from 10° C to 20° C.

After returning to Skierniewice, the boxes with mushrooms were put and kept in a temperature in accordance with the experimental

schemes. To eliminate the influence of transport conditions, boxes containing unirradiated mushrooms were also transported to Łódź and back to Skierniewice.

The retardation of mushroom growth and ageing was evaluated by measuring the increase of cap diameter, stem elongation, degrees of cap opening, percentage of mushrooms with open caps, discoloration of cap surface and sporulation.

The discoloration of caps was evaluated optically and in some experiments by using Hunter's Apparatus.

The sporulation was determined by the following test; after the storage period the mushrooms were kept in a thermostat at 21°C.

In the experiments, the occurrence of bacterial and fungal diseases was expressed as the percentage of mushrooms infected.

Fresh weight losses were also determined.

The results were evaluated statistically; standard deviation and analysis of variance were calculated. The LSD values were established for the 50/0 confidence level.

The results are presented in the form of diagrams. The values are the average for the particular series.

RESULTS

Effectiveness of irradiation with 2.0 kGy at various storage temperatures

Data presented in Figure 1 show that at a low storage temperature, 0-4°C, there was no effect of irradiation with 2.0 kGy on mushroom growth — cap size and stem elongation. The low temperature retarded mushroom growth effectively during 10 days of storage. The only difference observed was the better quality of the irradiated mushrooms at any evaluation time. After 10 days of storage the irradiated mushrooms were whiter and a higher percentage had closed caps.

The ionizing radiation retarded mushroom growth at the higher storage temperatures, $9-11^{\circ}C$ and $18-19^{\circ}C$.

At the temperature 9-11°C, the irradiated mushrooms grew slower than unirradiated ones. Their cap diameters increased a bit only until the fourth day of storage (Fig. 1A). Later, they remained the same size from the fourth to eighth day of storage. The caps of unirradiated mushrooms increased until the eighth day of storage.

The stem elongation of the irradiated mushrooms stored at 9-11°C was also retarded in comparison with unirradiated ones (Fig. 1B); the differences are not statistically significant.

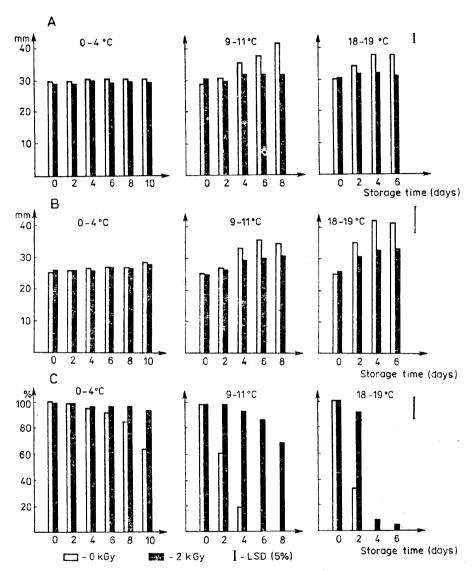


Fig. 1. Effect of ionizing radiation on retardation of mushroom growth at different storage temperatures; A — cap diameter. B — stem length, C — mushrooms with closed caps

The greatest differences between irradiated and unirradiated samples occurred in the cap opening of mushrooms at any time of evaluation (Fig. 1C). On the 4th storage day at $9-11^{\circ}$ C about $90^{\circ}/_{0}$ of irradiated mushrooms had closed caps, while at that time less than $20^{\circ}/_{0}$ of the unirradiated mushrooms had closed caps. After 6 storage days the caps of $80^{\circ}/_{0}$ of irradiated mushrooms remained closed, while caps of unirra-

diated mushrooms were all open. The color and the other quality properties as well as lack of diseases received better evaluation in the case of irradiated mushrooms than unirradiated ones. After 8 days of storage the irradiated mushrooms were marketable, while unirradiated were not edible.

At a storage temperature of 18-19°C the differences between the irradiated and unirradiated mushrooms occurred earlier (Fig. 1), after only 2 storage days. The differences are especially great when the influence of irradiation on cap opening is compared. After 2 storage days at 18-19°C, about 90% of irradiated mushrooms had closed caps, while only about 30% of the unirradiated mushrooms had closed caps. The unirradiated mushrooms stored at 18-19°C for 6 days deteriorated completely and were useless; they were over-grown and discolored, had dark gills and spreading spores, and were covered with molds and spots of bacterial diseases. At the same time the irradiated mushrooms were of much better quality, less grown and whiter; infection with diseases occurred in only 10% of the mushrooms.

The comparison of cap surface discoloration of irradiated and unirradiated mushrooms made on the basis of measurements with Hunter's Apparatus showed significant differences in favor of irradiated mushrooms.

Fresh weight losses during storage at various temperatures are shown in Figure 2. The irradiation did not influence the losses; they were affected mainly by storage temperature and increased significantly with

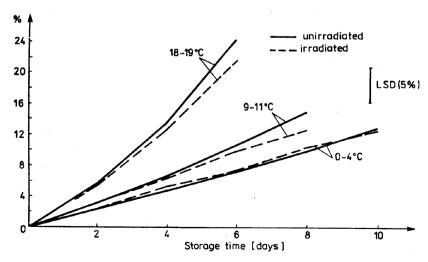


Fig. 2. Fresh weight losses of irradiated and unirradiated mushrooms stored at different temperatures

increasing storage temperature. After 6 days of storage at 18-19°C the losses were twice as great as at 9-11°C.

The results of further experiments (Fig. 3A, B, C) confirmed the effectiveness of the irradiation dose of 2.0 kGy on the retardation of mushroom growth and quality preservation for storage temperatures 10°C and 16°C. It can be stated that post-harvest mushroom growth occurred until the 5th day; later it stopped. The irradiation retarded stem elongation more than cap size increase (Fig. 3A, B). Stem growth of unirradiated mushrooms was somewhat lower at 16°C than at 10°C.

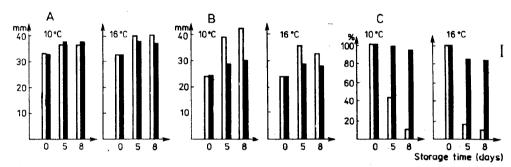


Fig. 3. Effect of ionizing radiation (2.0 kGy) and storage temperatures on retardation of mushroom growth; A — cap diameter, B — mushroom height, C — mushrooms with closed caps

Similarly as in previous experiments, the dose of 2.0 kGy affected significantly cap opening at both temperatures (Fig. 3C). After 8 days of storage at 10° C, about 90^{0} /₀ of irradiated mushrooms had their caps closed, while unirradiated ones had only about 10^{0} /₀. The data for storage at 16° C are 80^{0} /₀ and 10^{0} /₀, respectively.

The occurrence of fungal and bacterial diseases (Fig. 4A, B) was markedly decreased by irradiation and strongly depended on storage temperatue, particularly in the case of fungal diseases. After 8 days of storage at 16° C, the unirradiated mushrooms were all moldy while only $10^{0}/_{0}$ of the irradiated mushrooms were moldy. The dominant fungal disease was identified as *Acremonium album*. At a storage temperature of 10° C, visible infection with fungal diseases was lower.

The occurrence of bacterial diseases was also affected by irradiation. The differences between the irradiated and unirradiated mushrooms were significant (Fig. 4B). The dominant bacterial disease was brown spots (*Pseudomonas tolaasi*). After 8 days of storage at 10°C, less than 20% of the irradiated mushrooms were covered with brown spots, while about 70% of the unirradiated ones were covered with them. When stored at

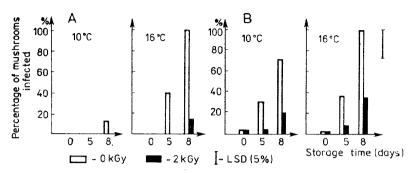


Fig. 4. Effect of ionizing radiation (2.0 kGy) and storage temperatures on the infection of mushrooms caused by fungal (A) and bacterial (B) diseases

 16° C, 30° /o of the irradiated mushrooms were infected compared with 100° /o of the unirradiated mushrooms.

The measurements of fresh weight losses confirmed that the irradiation had no effect on them. Similary as in previous experiments, the losses depended on storage temperature. After 8 days of storage at 16° C the fresh weight losses were twice as great as at 10° C.

Effectiveness of lower radiation doses

The results of the experiments with the doses of 0.5 kGy and 1.0 kGy carried out at storage temperature 10°C are given in Figures 5, 6 and 7. Irradiation did not affect the cap growth (cap diameter increase, Fig. 5A).

Visible differences, mostly statistically significant, occurred in the increase of elongation (Fig. 5B, C). The dose effect can be seen. The dose effect was also seen upon the examination of the influence of irradiation on cap opening (Fig. 6). Generally, the dose of 1.0 kGy gave similar

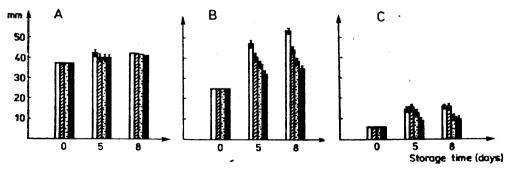


Fig. 5. Effect of various doses of ionizing radiation (0, 0.5, 1.0, 2.0 kGy) on retardation of mushroom growth at storage temperature 10°C; A — cap diameter, B — mushroom height, C — stem length

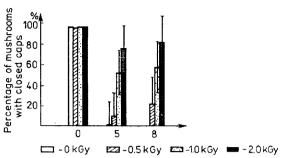


Fig. 6. Influence of various doses of ionizing radiation on retardation of mushroom cap opening

effects to the dose of 2.0 kGy. The dose of 0.5 kGy was less effective.

The quality properties, color and texture, showed the dose effect too; higher doses increased retention of quality.

The test for sporulation of mushrooms indicated that all applied doses retarded it. No spore spreading was observed in irradiated mushrooms, even at the low dose of $0.5~\mathrm{kGy}$; unirradiated mushrooms readily spread off spores.

The occurrence of fungal disease (*Acremonium album*) and bacterial brown spots (*Pseudomonas tolaasi*) was strongly affected by irradiation (Fig. 7) and the dose effect was seen.

After 8 days of storage at 10°C, 50% of the unirradiated mushrooms

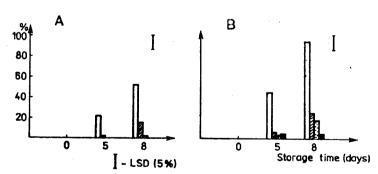


Fig. 7. Influence of various doses of ionizing radiation on the occurrence of mushroom diseases; A — fungal diseases, B — bacterial diseases

were covered with white mold (Acremonium album); only 10% of the mushrooms irradiated with the dose 0.5 kGy were infected. At higher doses, 1.0 kGy and 2.0 kGy, molding did not occur. All applied doses affected the occurrence of bacterial brown spots (Pseudomonas tolaasi). After 8 days of storage all the unirradiated mushrooms were covered

with brown spots, while only $20^{\circ}/_{\circ}$ of the mushrooms irradiated with the dose of 0.5 kGy were infected. Higher doses decreased the extent of bacterial infection; a low percentage of the mushrooms was infected.

DISCUSSION

The results presented confirm the fact that irradiation of mushrooms retards their growth and ageing. The effect is strongly dependent on the dose and thermal conditions of storage after irradiation. As a consequence of the retarded growth and maturity, the quality properties of mushrooms may be retained for a longer time, enabling the prolongation of storage time. In our experimental and storage conditions the dose of 2.0 kGy, applied within a few hours after harvest, on mushrooms at the "cup closed" stage and at a storage temperature of 10°C (at high humidity), caused an increase of shelf life to 8 days; similar storage time prolongation was obtained at 16°C. At the storage temperature of 18-19°C, the prolongation of the shelf-life of irradiated mushrooms was limited to 6 days.

At the low storage temperature, $0\text{-}4^{\circ}\text{C}$, an effect of irradiation with 2.0 kGy on mushroom growth was not observed until 10 days of storage. This temperature retarded mushroom growth well. Only the quality of irradiated mushrooms was better in comparison with unirradiated.

Similar results were obtained in experiments carried out by a number of investigators. Some differences might be caused by different experimental conditions (mushrooms state, storage parameters, and irradiation technique).

The results of our experiments are similar to those obtained by Langerak (1971). He stated that irradiation with 2.0 kGy and storage at 3-5°C increased the shelf-life by 10 days; storage at 8-10°C by 7 days, and at 20°C by 5 days. In experiments by Farkas (1975) irradiation with 3.0 kGy and storage at 16-18°C (at 65% RH) increased the shelf-life to 3 days. Kovaćs and Vas (1974) increased the shelf-life to 13 days by irradiation with 2.0 kGy and storage at 6-8°C (65% RH); at a higher temperature, 16-18°C, the shelf-life increased to 11 days. In an earlier investigation by Staden (1965), mushrooms irradiated with 2.0 kGy, stored at 10°C, kept closed caps for 12 days; the unirradiated mushrooms were all open after 7 days. Markakis (1968) obtained similar results with the same dose when stored at 5°C for 12 days. Salkova et al. (1978) reported that mushrooms irradiated with a dose of 2.0 kGy and stored at 2°C kept their quality for 20 days; at a storage temperature of 15°C, quality was maintained for 7 days. But

at the storage temperature of 24°C, an effect of irradiation did not occur; irradiated mushrooms deteriorated as quickly as unirradiated ones.

Irradiation slowed the darkening processes, probably through the retardation of ageing. In all experiments we observed that irradiated mushrooms kept their whiteness for a longer period and at any evaluation time the difference was visible and verifiable by measurements using Hunter's Apparatus.

As a consequence of retarded growth, the irradiated mushrooms also kept their texture for a longer time. The influence of irradiation on maintaining the color was also reported by Staden (1965), Gill et al. (1969), Langerak (1971), Kovaćs and Vas (1974 A, B), Barkai-Golan and Padova (1981). Some of these authors also demonstrated better texture of irradiated mushrooms after storage in comparison with unirradiated ones. They found no differences in flavor and taste between treated and untreated mushrooms.

In the investigation of the effectiveness of lower doses of irradiation we observed that the dose of 1.0 kGy gave results approaching the results obtained with a dose of 2.0 kGy when the storage temperature was 10°C. The dose of 0.5 kGy was less effective. The dose effect was observed on the growth parameters: mushroom height and cap opening.

Staden (1965) looking for the most effective low radiation dose investigated a range of doses from 0.5 kGy to 2.5 kGy. The influence on cap opening was the same at 1.0 kGy and 2.0 kGy. Markakis (1968) stated that doses of 0.25, 0.5 and 1.0 kGy similarly affected cap opening. Gill et al. (1969) found that low doses 0.1-0.5 kGy, were less effective; these doses influenced cap opening more strongly than stem elongation. The latter was affected by dose of 1.0 kGy and higher. Markakis (1968), Kovaćs and Vas (1974a, b) concluded that for unripe mushrooms, irradiated shortly after picking, the low dose (0.25 kGy) retarded mushroom growth effectively. According to Farkas (1975) the dose 0.25 kGy was effective for storage at temperature 5-8°C at 70-90% RH. Salkova et al (1978) regarded the doses 1.5-3.0 kGy as most effective. Barkai-Golan and Padova (1981) stated that the dose of 0.5 kGy and higher inhibited cap opening, stalk elongation at storage temperature 15°C.

In our experiments all applied doses (0.5, 1.0 and 2.0 kGy) retarded the sporulation of mushrooms. Ionizing radiation may influence this phenomenon in various ways. Kovaćs (1982 a, b) stated that irradiation slowed down the ageing processes, the cytokinin activities as well as sporulation. As the result of irradiation, the spore count decreased as a function of the dose; after a dose of 2.0 kGy and 5-days storage, the spore count did not exceed the original value, while in non-treated

mushrooms the spore count increased five-fold. Similarly, spores of treated mushrooms maintained their original color while non-treated ones became entirely dark.

Another radiation effect indicated in our experiments worth stressing is that irradiation strongly reduced the incidence of fungal and bacterial diseases. This fact might be of importance in storage problems and deserves more attention.

It was already reported by Staden (1965, 1966), that irradiated mushrooms remained mold-free during storage. Gill et al. (1969) also observed less mold growth on irradiated mushrooms. Salkova et al. (1978) showed the decrease in pathogenic fungal spores of artificially infected mushrooms after irradiation with 2.0 and 3.0 kGy.

As a consequence of mushroom growth retardation and significant control the occurrence of fungal and bacterial diseases, in our experiments irradiation with a dose of 2.0 kGy allowed the self-life of mushrooms to increase to 8 days at storage temperature of 10°C and 16°C, and to 6 days at storage temperature of 18-19°C.

The practical application of the irradiation method in mushroom storage may be of value in the case when cold storage (near 0° C) would be more expensive than the treatments with ionizing radiation and storage at 10° C and even higher temperatures till 16° C. These temperatures are available in common storage rooms, ventilated with outside air, during colder seasons.

Acknowledgment

The authors are grateful to the Department of Microbiology of the Research Institute of Vegetable Crops in Skierniewice for help in the identification of fungal diseases of mushrooms.

We also thank Miss M. Walczak and Ing. G. Świderski for technical assistance during performence of the experiments.

REFERENCES

- Barkai-Golan R., Padova R., 1981. Extending the shelf-life of cultivated mushrooms by gamma irradiation. Hort. Abst. 51, 3 (poz. 1718).
- Farkas J., 1975. Present status and prospects for the commercialization in Hungary of irradiated food items for human consumption. Requirements for the Irradiation of Food on a Commercial Scale. Proc. of Panel, Vienna 1975. IAEA, op. 37-57.
- Gill W. I., Nicolas R. C., Markakis P., 1969. Irradiation of cultured mushrooms. Food Technology 23: 385-388.
- Kovaćs E., Vas K., 1974a. Effect of ionizing radiation on some organoleptic characteristics of edible mushroom. Acta Alimentaria 3: 11-17.

- Kovaćs E., Vas K., 1974b. Effect of ionizing radiation on post-harvest ripening processes of cultured mushrooms (*Agaricus bisporus*) with special references to the rates of respiration and of ethylene production. Acta Alimentaria 3: 19-25.
- Kovaćs E., 1982a. Cytokinin activity of *Agaricus bisporus*. Acta Alimentaria 11: 169-178.
- Kovaćs E., 1982b. Combined effect of kinetin and radiation treatment on the cap opening of *Agaricus bisporus*. Acta Alimetaria 11: 179-187.
- Langerak D. J., 1971. The influence of irradiation and packaging upon the keeping quality of fresh mushrooms. Proc. Mushrooms Science 8: 221-230.
- Markakis P., 1968. Irradiation of mushrooms. Mushrooms News 17: 8-11.
- Salkova Z., Kubin K., Stanek M., Horaćek P., 1978. Present status of food irradiation in Czechoslovakia. Proc. of Symp. Food Preservation by Irradiation, Wageningen IAEA 1: 43-51.
- Staden O. L., 1965. Radiation preservation of fresh mushrooms. Mushroom Science 6: 457-461.
- Staden O. L., 1966. Experiences with the irradiation of vegetables in the Netherlands. Proc. of Symp. Food Irradiation, IAEA, pp. 609-617.
- Śmierzchalska K., Gubrynowicz E., 1979. Stosowanie promieniowania jonizującego w przechowalnictwie pieczarek. Post. Nauk Rol. 2: 67-80.

Wpływ promieniowania jonizującego i temperatury przechowywania na wzrost i jakość pieczarek (Agricus bisporus)

Streszczenie

W latach 1979-1981 przeprowadzono doświadczenia ścisłe nad wpływem promieniowania jonizującego dawki 2,0 kGy i różnych temperatur przechowywania grzybów po napromieniowaniu na hamowanie procesów wzrostu i starzenia się grzybów. Badano również skuteczność mniejszych dawek 0,5 i 1,0 kGy.

Świeżo zebrane pieczarki dobrej jakości napromieniowywano w terminie 5-6 godzin po zbiorze, dawką 2,0 kGy, następnie umieszczono w różnych temperaturach. Efekt zastosowanej dawki zależał w dużym stopniu od temperatury przechowywania grzybów. W temperaturze 0-4°C był najmniej widoczny. Taka temperatura hamowała wzrost pieczarek zupełnie dobrze w ciągu 10 dni. Jednakże grzyby napromieniowane charakteryzowały się lepszą jakością, były bielsze i większy procent miał zamknięte kapelusze.

W wyższych temperaturach przechowywania pieczarek wpływ napromieniowania wystąpił wyraźniej. Dawka 2,0 kGy przy przechowywaniu w temperaturze 10°C hamowała wzrost elongacyjny, przyrosty średnicy kapeluszy i ich otwieranie, a także ciemnienie i zarodnikowanie. Zmniejszała również stopień porażenia pieczarek chorobą bakteryjną (Pseudomonas tolaasi) i grzybową (Acremonium album).

Podobne wyniki uzyskano przy przechowywaniu w temperaturze 16°C.

W wyższych temperaturach 18-19°C efekt napromieniowania dawką 2,0 kGy był również widoczny, z tym, że pieczarki znacznie szybciej ulegały niekorzystnym zmianom.

Poprzez zahamowanie procesów wzrostu pieczarek i rozwoju chorób bakteryjnych i pleśni, napromieniowanie dawką 2,0 kGy przedłużało okres składowania do 8 dni, gdy pieczarki przechowywano w temperaturze 10-16°C, i do 6 dni, gdy

przechowywano je w temperaturze 18-19°C. W tym czasie pieczarki nienapromieniowane były bardzo złej jakości i nie nadawały się do spożycia.

Mniejsza dawka promieniowania jonizującego 1,0 kGy i przechowywanie w temperaturze 10°C, wykazywała działanie zbliżone do dawki 2,0 kGy. Dawka 0,5 kGy była mniej skuteczna. Wszystkie stosowane dawki promieniowania jonizującego hamowały osypywanie zarodników.

Praktyczne zastosowanie metody radiacyjnej w przechowywaniu pieczarek może mieć znaczenie, gdy przechowywanie w niskich temperaturach okaże się kosztowniejsze niż napromieniowanie grzybów i przechowywanie w temperaturze 10°C, a nawet wyższych. Takie temperatury panują u nas w zwykłych przechowalniach w ciągu chłodniejszych miesięcy wiosny i jesieni.