

Irradiated samples indicate only a small increase of the PPSL signal, whereas with unirradiated ones the increase of the signal is significant.

Application and limitations of measures

The method of detection of irradiated food by means of PPSL has been positively tested in inter-laboratory tests for samples of shellfish (*e.g.* prawns), herbs, spices and seasoning [9-11].

PPSL sensitivity depends on the quantity and type of minerals present in the individual sample. Signals of the intensity below the lower threshold (T_1) are generally associated with unirradiated material, but sometimes can be also derived from low sensitivity irradiated materials. In general, calibrated PPSL measurements are recommended for shellfish with low mineral contents and „clean” spices (*e.g.* nutmeg, white and black pepper) to avoid false negative results [9,14]. According to our experience, for the examination of any sample delivered from our clients the calibrated PPSL measurement should be always adapted.

Multicomponent food products like curry powder, for example, and blended seasonings may contain the debris of minerals of low PPSL sensitivities, in which case calibrated PPSL may also provide unclear results. In such a case it is necessary to turn to TL measurements.

Food products classified in the course of our investigation as such that may provide unclear results of PPSL measurements are: garlic powder, carrot pepper (leaves), sweet pepper (powder), black pepper (grains), black pepper (ground), clove (whole), dried dill (powder).

The presence of salt in a product given for examination intensifies so much the PPSL signal intensity that its contribution dominates to an extent which masks effectively signals from any irradiated ingredient. The dominance of the luminescence from crystalline salts in a product makes the signals from irradiated components undetectable. An admixture to a product of the following salts makes the examination of by PPSL method not rational: sodium chloride (domestic salt), natrium sorbitan, sodium benzoate, monosodium glutamate, Arabic gum.

It has to be strongly stressed that the examination of samples containing the above ingredients may also cause the damage of photomultiplier and is prohibited.

Sometimes hydration of a product leading to full dilution of salt and its elimination followed by drying and PPSL measurement can both identify and rectify this situation.

Conclusions

The PPSL method can be successfully used for the detection of irradiation in pure spices, herbs

and seasonings as well as in most of multicomponent blends of spices, herbs and seasonings [6,14,15].

Screening by means of the PPSL apparatus is easy, effective and first and above all inexpensive. The method provides the fastest way to gain final results whether food product is irradiated. By comparison with the TL method, preparation of samples is simple, much quicker and takes not longer than one hour instead of few days by the TL method. However, in ambiguous results of PPSL, the validated TL method should be always used [16].

References

- [1]. Pinnioja S., Siitari-Kauppi M., Jernström J., Lindberg A.: *Radiat. Phys. Chem.*, **55**, 743-747 (1999).
- [2]. Sanderson D.C.W., Slater C., Cairns K.J.: *Radiat. Phys. Chem.*, **34**, 915-924 (1989).
- [3]. Soika Ch., Delincée H.: *Lebensm.-Wiss. Technol.*, **33**, 440-443 (2000), in German.
- [4]. The SURRC Pulsed Photostimulated Luminescence (PPSL) Irradiated Food Screening System. Users Manual. Royal Society of Chemistry, Cambridge 2004, 17 p.
- [5]. Bluszcz A.: *Zeszyty Naukowe Politechniki Śląskiej*, **86(1434)**, 11-17, 25-47 (2000), in Polish.
- [6]. EN 1788:2001: Foodstuffs – Thermoluminescence detection of irradiated food from which silicate minerals can be isolated.
- [7]. Directive 1999/3/EC of the European Parliament and of the Council of 22 February 1999 on the establishment of a Community list of food and food ingredients treated with ionising radiation. *Off. J. European Communities L 66/24-25* (13.3.1999).
- [8]. CEN/TC 275/WG 8 N 127: Detection of irradiated food using photostimulated luminescence. 1999.
- [9]. Detection of irradiated samples. European Patent No. 0 699 299 B1.
- [10]. PN-EN 13751:2003 (U): Artykuły żywnościowe – Wykrywanie napromieniowania żywności za pomocą fotoluminescencji.
- [11]. Sanderson D.C.W., Carmichael L., Fisk S.: *Food Sci. Technol. Today*, **12(2)**, 97-102 (1998).
- [12]. Sanderson D.C.W., Carmichael L.A., Naylor J.D.: *Food Sci. Technol. Today*, **9(3)**, 150-154 (1995).
- [13]. Sanderson D.C.W., Carmichael L.A., Naylor J.D.: Recent advances in thermoluminescence and photostimulated luminescence detection methods for irradiated foods. In: *Detection methods for irradiated foods – current status*. Royal Society of Chemistry, Cambridge 1996, pp.124-138.
- [14]. Huntley D.J., Godfrey Smith D.I., Thewald M.L.W.: *Nature*, **313**, 105-107 (1985).
- [15]. Sanderson D.C.W.: Detection of irradiated samples. Great Britain Patent No. 93-8542 GB 9308542.
- [16]. Guzik G.P., Stachowicz W.: *Pomiar luminescencji stymulowanej światłem, szybka metoda identyfikacji napromieniowania żywności*. Instytut Chemii i Techniki Jądrowej, Warszawa 2005, 16 s. Raporty ICHTJ. Seria B nr 3/2005 (in Polish).

DETECTION OF IRRADIATION IN CUTICLES OF COMMERCIAL SHRIMPS

Katarzyna Lehner, Waclaw Stachowicz

The detection of stable EPR (electron paramagnetic resonance) signal produced by the action of

ionising radiation in crustacea has been reported by several authors elsewhere [1,2]. The signal ob-

served was identical with that used for detection of irradiation in bone and eggshell by the EPR method. This signal is specific for hydroxyapatite that appears in some extent in exoskeletons of shrimps and crabs. However, the results of model studies on these products were not reliable enough. For that reason, crustacea are not quoted until now in European Standard EN 1786 among food products that can be examined by EPR method to prove their irradiation [3]. The most reliable results have been obtained with lobsters [4]. The results of the examination of various crustacea species indicated that the EPR signal is more or less influenced by the variety, origin and age of crabs.

Nowadays, the Laboratory for Detection of Irradiated Food has more and more orders for detection of irradiation in shrimps. Part of them are delivered in cuticles. A method adapted to accomplish the detection of irradiation in shrimps is a thermoluminescence measure. However, the analytical procedure is much time-consuming and needs several days to receive the final result.

The aim of present study was to prove, whether the EPR examination of cuticles taken from shrimps could be used as a screening method for the detection of irradiation in this product. The earlier results obtained with irradiated cuticles of shrimps only seem promising [5].

In a model study commercial shrimps were irradiated in a ^{60}Co source with doses of 1, 3 and 7 kGy. The technological dose recommended for microbial decontamination of shrimps is between 3 and 7 kGy.

Cuticles taken from the shrimp body were cleaned, dried and subsequently crushed to small pieces to be measured by the EPR method. The resultant spectra recorded with cuticles irradiated with 3 and 7 kGy are shown in Fig.1. The positions of coefficient g (spectroscopic splitting factor), specific for irradiated hydroxyapatite, are marked with arrows. The signal is a singlet of axial symmetry with $g_x=2.0035$, $g_y=1.9973$ and $g_z=2.0017$, respectively.

The positions of all three g 's in the magnetic field are easily distinguished in the spectrum of cuticle irradiated with 7 kGy (Fig.1b). However, in the spectrum of cuticle irradiated with 3 kGy the positions of g_z and g_x are not very well defined although the experienced EPR operator can establish them with a precision which could be perhaps

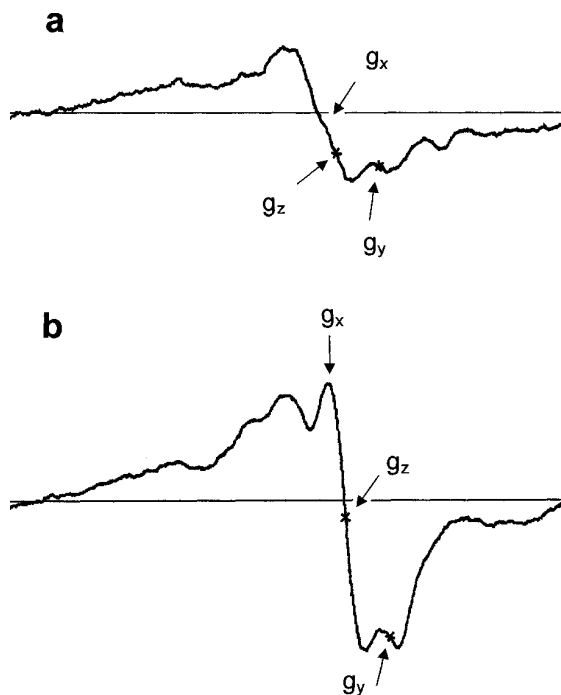


Fig.1. The EPR spectra (first derivatives) recorded with cuticles taken from shrimps irradiated with 3 kGy (a) and 7 kGy (b). The positions of g values specific for irradiated hydroxyapatite are marked with arrows: $g_x=2.0035$, $g_y=1.9973$, $g_z=2.0017$.

satisfactory enough for identification of radiation treatment (Fig.1a).

In conclusion, it can be postulated that the EPR measurement of cuticles of shrimps can be adapted in the Laboratory as preliminary, screening test proving the irradiation of shrimp. It has to be stressed, however, that the lack of a specific, hydroxyapatite born EPR signal in the spectrum cannot suggest that sample was not irradiated.

References

- [1]. Desrosiers M.F.: *J. Agric. Food Chem.*, **37**, 96-100 (1989).
- [2]. Raffi J.J., Agnel J.P.: *Sciences des Aliments*, **10**, 387-391 (1990).
- [3]. European Standard EN 1786:1996: Foodstuffs – Detection of food containing bone. Method by ESR spectroscopy.
- [4]. Stewart E.M, Stevenson M.H., Gray E.: *Appl. Radiat. Isot.*, **44**, 1-2, 433-437 (1993).
- [5]. Morehouse K.M., Desrosiers M.F.: *Appl. Radiat. Isot.*, **44**, 1-2, 429-432 (1993).

DSC STUDIES OF RETROGRADATION AND AMYLOSE-LIPID TRANSITION TAKING PLACE IN GAMMA-IRRADIATED WHEAT STARCH

Krystyna Cieřla, Ann-Charlotte Eliasson^{1/}, Wojciech Gluszewski

^{1/} Department of Food Technology Engineering and Nutrition, University of Lund, Sweden

The course of gelatinisation and retrogradation occurring during heating of starch and flour suspensions depend on the structure of starch granules. In the case of wheat flour, retrogradation depends additionally on the presence of lipids. In fact,

binding of lipids to the polysaccharide chains was found to resist recrystallisation of starch gels.

Our previous studies have shown that degradation resulting from gamma irradiation induces a decrease in order of starch granules [1,2] and