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## Comparison of Nutritional Composition and Histological Changes of the Soybean Seeds Cultivated by Conventional and Organic Farming Systems after Long-Term Storage – Preliminary Study –

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We investigated the effect of hermetic long-term (9-year) storage at room temperature on the chemical compositions and histological structure of soybean seeds that had been cultivated by two farming systems [conventional farming (CF) or organic (natural) farming (OF)]. Soybean seeds had been harvested from plants cultivated in the field under CF practices, or in a neighboring field where OF practices had been used for over 3 years. Protein content, calcium (Ca), potassium (K) and phosphorus (P) content of OF soybean seeds were higher than CF seeds just after harvesting. However, after long-term storage, protein content of OF soybean seeds was significantly higher than CF seeds. In addition, there were many minerals with a higher content (in particular, K) of OF soybean seeds. The breaking strength and pH of OF soybean seeds were higher than CF ones. Optical microscopic examination revealed collapsed tissue and gaps in the internal and external parts of CF soybean seeds. Transmission electron microscopic photographs of CF soybean seeds confirmed the presence of collapsed cell walls and protein bodies. These results indicate that the type of farming system (inorganic or organic fertilizers) used results in differences in chemical compositions and histological structure of product during storage. In addition, our results indicated that high Ca content of OF soybean seeds strengthened cell wall structure, and prevented the loss of contents during storage.

KEYWORDS: histology, soybean seed, conventional farming, organic farming, storage

### INTRODUCTION

Various methods, such as cool storage or CO<sub>2</sub> gas storage, are used to keep farm products such as grains and seeds (Calderwood *et al.*, 1984; Zhou *et al.*, 2002). However, the quality of the products inevitably deteriorates during storage (Hoshi *et al.*, 1982; Sripichitt *et al.*, 1989; Yamamoto and Shirakawa, 1999; Matsue *et al.*, 2003; Hou and Chang, 2004a, b). A number of physiological and physicochemical changes take place and are usually termed ageing (Narayan *et al.*, 1988a, b; Zhou *et al.*, 2002). These changes affect colour, flavour, texture and the occurrence of mold (Wilkins and Scholl, 1989; Zhou *et al.*, 2002). Until now, there have been few reports (Benge *et al.*, 2000; Asami *et al.*, 2003) about changes in quality (especially chemical composition) of organically grown and conventional products under storage; furthermore, reports of comparative tests have shown that quality after storage is better in organic products than in conventional products (Asami *et al.*, 2003; Lotter, 2003). This suggests that these differences may result from differences in the farming systems, in particular, the form of fertilizer (organic versus

inorganic) used as a source of nutrition. However, there is little data with regard to the chemical composition and histological changes of products from conventional farming (CF) or organic (natural) farming (OF) systems after long-term storage. We assume that the degree of deterioration is determined, to some extent, by the histological structure of the cell walls and chemical composition (Narayan *et al.*, 1988a,b; Zhou *et al.*, 2002; Asami *et al.*, 2003).

The method of natural farming on equal terms with OF was advocated in Japan by Okada (1982), who emphasized the importance of soil health and microbial biodiversity and is conceptually similar to the suggestions of France (1921). Taken together, our results and the reports by Okada (1982) and France (1921) indicate that the type of farming system affects the morphology of the products, and that the effect was mainly due to the type of fertilizer that was used, namely chemical or organic (Nakamura, 1988, 1989; Nakamura *et al.*, 2003). However, the effect of the type of farming on the morphogenesis of products or change during cooking and storage has not been clarified. On the other hand, the OF system has attracted attention throughout the world (Reganold *et al.*, 2001; Tilman *et al.*, 2002; Altieri and Nicholls, 2003). However, until now the chemical and histological differences between CF and OF products have not been delineated (Woese *et al.*, 1997; Brandt and Mølgaard, 2001). In Japanese agriculture, use of the OF system is presently in a minority, and the volume of the products produced is small (Neera *et al.*, 1992, 1999).

We studied the chemical compositions and histologi-

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cal changes of soybean seeds that had been cultivated by CF or OF practices followed by harvest and long-term storage to evaluate the significance of organic substances and soil biota on the chemical compositions and cell morphology of soybean seeds.

## MATERIALS AND METHODS

### Materials

Seeds of soybean (*Glycine max* L. Merr. variety Horei) were sampled during the harvest season (in Nov.) from upland fields of CF and OF in the Tohoku National Agricultural Research Center in Fukushima city in a district in northeastern Japan.

Historically, the CF fields had been farmed conventionally (receiving chemical fertilizers and agricultural chemicals that were standard for the respective district), while the OF fields had been cultivated naturally for over 3 years without chemical fertilizers or agricultural chemicals and using no-till planting and weeding (which had been performed several times during the course of this study). The soil conditions and agricultural practices in Fukushima were reported in detail by Nakamura *et al.* (2003). Approximately 500 g of each material was used for this study.

### Storage conditions

The materials were soybean seeds that had been harvested in 1995 and stored for 9 years under hermetic conditions at room temperature and shield from light.

### Measurement of dry matter, protein, lipid and ash contents after harvest and long-term storage

Samples (100 g) from harvest and long-term storage (in Nov. 2004) were powdered, and the proximate composition was measured using the methods of Anonymous (1990). Moisture content was measured by heating the sample to 110 °C for 18 h. Crude protein content was measured by the Kjeldahl method. The coefficient of conversion from the amount of nitrogen to crude protein content was 5.71 based on the Standard Tables of Food Composition in Japan (STFCJ 2001). Crude lipid content was measured by the Soxhlet Extraction System (Sanyoh MFG Co. Ltd., Osaka, Japan). Crude ash content was measured by heating the sample to 600 °C for 24 h.

Samples after harvest were measured for protein and lipid content only. In addition, each food product type was assayed in one (after harvest) or ten (after long-term storage) replicates.

### Simple qualitative analysis of mineral compositions after harvest and long-term storage

To prevent trace metal contamination of the samples by laboratory equipment, all laboratory ware was soaked in 2 M HNO<sub>3</sub> for 48 h, rinsed five times with distilled water, and then five times with deionized water prior to use. Deionized water, of resistivity greater than 18 /cm and pressure of 4 kg/cm<sup>2</sup> at 25 °C, was prepared by passing singly ultra-pure water through Ultra Pure

Water System (type: CPW-100-CCS-1) (Advantec Toyo Kaisha, Ltd., Tokyo, Japan). For measurement of toxic metals, tissue samples (3 g) were digested with concentrated nitric acid (Wako Pure Chemical Industries, Ltd., Osaka, Japan). Samples were heat-treated at 120 °C on a hot plate with hydrogen peroxide (Wako Pure Chemical Industries, Ltd., Osaka, Japan) until sample solution became transparent. By adding ultra-pure water, sample solutions were adjusted to 100 mL total volume.

Mineral content of each sample was analyzed for detectable minerals by an Inductively Coupled Plasma – Atomic Emission Spectrometer (IRIS intrepid XDL) (Thermo Electron Co., Waltham, MA, USA). The final reported values are the mean of three replicate determinations. However, data of sodium (Na) was not shown because a glass flask for the heat-treatment. Analysis conditions were: delay time, 0; sample substitution time, 60; and largest integral time, 20 (low) and 10 (high).

Samples after harvest were measured calcium (Ca), potassium (K) and phosphorus (P) levels only. In addition, each sample was assayed in one (after harvest) or three (after long-term storage) replicates.

### Measurement of the breaking strength after long-term storage

Ten soybean seeds of each group, which had absorbed water for 24 h at room temperature, were divided into half. A blade plunger was used to cut the middle of seeds at a speed of 2 cm/min, and maximum force was recorded by a rheometer (RT-1002A, Fudoh, Tokyo, Japan) 20 times for each group and was regarded as the breaking strength.

### Measurement of pH after long-term storage

The pH of soybean seeds after long-term storage was measured using by a pH meter (F-53; Horiba, Kyoto, Japan). The soybean samples (5 g) were combined with distilled water (20 mL) in a test tube, and were allowed to absorb the water at 4 °C for 12 h. Samples were homogenized for 1 min at 10,000 rpm at low temperature. The pH of the mixture was directly measured by the electrode of a pH meter. Each sample was assayed in three replicates.

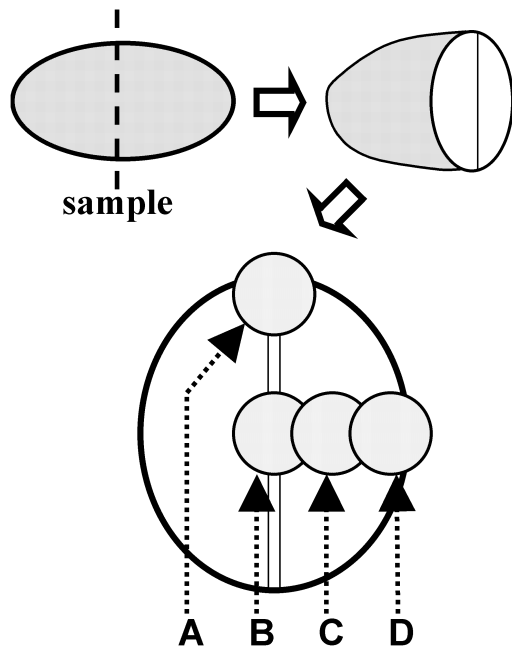
### Scanning electron microscopy after long-term storage

The soybean seeds were cut 1 × 1 mm of surface of seed coats and surface of seeds which stripped off seed coat, and then mounted on aluminium holders with the surface of seed coats and surface of the seeds which stripped off seed coat. The specimens were coated with Pt-Pd by an Ion Sputtering Device (Quick Auto Coater JFC-1500, JEOL). The surface of seed coats and surface of seeds which stripped off seed coat were examined under a scanning electron microscope (S-2380N, Hitachi, Tokyo, Japan).

### Optical microscopy after long-term storage

In Nov. 2004, each sample was analyzed by optical

microscopy. Ten equal-length soybean seeds were randomly selected from storage bottles of each product group (CF and OF); they were fixed in 5% glutaraldehyde (in 0.1 M phosphate buffer, pH 7.4), stored at 4 °C for 2 weeks, cut in two longitudinally, and stored in the fixative at 4 °C for 1 additional week. After dehydration through a series of 50–to–100% ethanol washes, samples were embedded in Technovit (Technovit 7100; Ohken Co., Tokyo, Japan), and 3  $\mu$ m-thick, longitudinal sections were sliced using a microtome (Nippon Optical Works, Tokyo, Japan). Sections were stained with a 0.1% toluidine blue and analyzed under an optical microscope (BX-50; Olympus, Tokyo, Japan) (**Fig. 1**).



**Fig. 1.** Diagrams of cross-sections of soybean seeds used for optical microscopy (OM).

#### Transmission electron microscopy after long-term storage

Ten same-length soybean seeds (1 × 1 × 5 mm) selected by random sampling from each product group (CF and OF) were fixed in 5% glutaraldehyde (in 0.1 M phosphate buffer, pH 7.4) for 1 week. After being washed with the sample buffer, they were post-fixed with 1% osmic acid medium in 0.1 M phosphate buffer (pH 7.4). Dehydration using a 50–to–100% ethanol series was performed before saturation with propylene oxide. After embedding the seeds in epoxy resin (Epon 812; Ohken Co., Tokyo, Japan), ultra-thin sections (90 nm thick) were prepared using an ultra-microtome (MT 6000 Sorvall Instruments; Du Pont Co., Delaware, USA). They were stained with 1% uranyl acetate and lead citrate and observed under a transmission electron microscope (H-800; Hitachi, Tokyo, Japan) with an accelerating voltage of 100 kV.

#### Statistical analysis

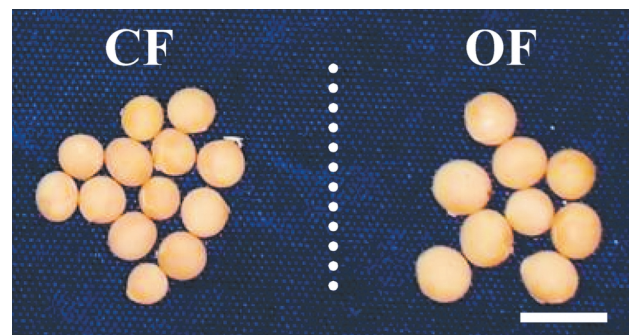
Means and standard errors of the proximate composition and proximate mineral compositions of each sam-

ple were calculated after long-term storage. A *t*-test was used to determine significant differences between the proximate compositions of CF and OF products after long-term storage.

## RESULTS AND DISCUSSION

Changes in the chemical and histological properties of farming products occur after harvest and during storage. For example, there are changes in smell, color and texture (Narayan *et al.*, 1988a, b; Zhou *et al.*, 2002). These changes are caused by consumption of organic compounds (such as starch, protein, lipids, vitamins, etc.) by respiratory processes and oxidation (Clark and Snyder, 1991; Tanteeratarm *et al.*, 1989; Angulo *et al.*, 1998; Zhou *et al.*, 2002; Rukunudin *et al.*, 2004). Furthermore these chemical changes are effected by the humidity (Patil *et al.*, 1986) and temperature conditions (Kaushik *et al.*, 1987; Sorour and Uchino, 2004) during storage, and they are accompanied by histological changes (especially in texture) (Ojijo *et al.*, 2000a, b; Zhou *et al.*, 2002).

The applied farming system itself affects the chemical and histological properties of the products (Yoshida *et al.*, 1984a, b, c; Bengé *et al.*, 2000; Brandt and Mølgaard, 2001). In the present study, we observed changes in the chemical compositions and histological structure of the CF and OF soybean seeds under long-term storage for 9 years under hermetic conditions at room temperature and shielded from light. In the present study, each of the sample products was kept in good condition, and there was no evidence of mold, harmful insect damage, or putrefaction (**Fig. 2**). There



**Fig. 2.** Photograph of each sample after long-term (9 years) storage. CF: conventional farming, OF: organic farming. Bar = 1 cm.

**Table 1.** Protein, lipid contents and mineral compositions of the soybean seeds by conventional (CF) and organic (OF) farming systems after harvest

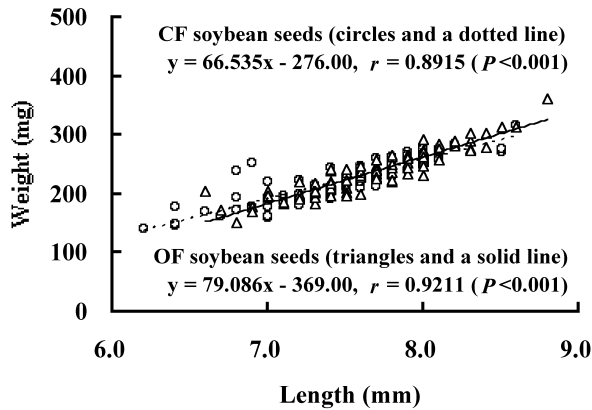
	Farming system	
	CF	OF
Protein (% in dry weight)	33.1	35.1
Lipid	18.1	17.2
Ca (mg · kg <sup>-1</sup> in dry weight)	1890	2010
K	17500	18100
P	4210	4510



**Table 2.** Protein, lipid and ash contents of the soybean seeds by conventional (CF) and organic (OF) farming systems after long-term storage

	Farming system				
	CF		OF		<i>t</i> -Test
	Mean	S.D.	Mean	S.D.	
Dry matter (%)	90.5	0.36	91.0	0.31	$P < 0.05$
Protein (% in dry weight)	32.9	0.64	36.2	1.20	$P < 0.001$
Lipid	8.71	1.07	8.75	0.77	$P > 0.05$
Ash	3.82	0.14	3.96	0.16	$P > 0.05$

S.D.: standard deviation.

**Fig. 3.** The correlation of length and weight of soybean seeds after long-term (9 years) storage.  $n = 100$ .**Table 3.** Difference of mineral contents of soybeans seeds by conventional (CF) and organic (OF) farming systems after long-term storage.

	Farming system	
	CF	OF
Difference		
> + 100 mg·kg <sup>-1</sup>	–	K ( + 4250 mg)
+ 50 ~ 100	–	Ca, P
+ 10 ~ 50	S	Mg
+ 1 ~ 10	Zn	B, Ba, Ga, Mn, Sm
Not difference		
	Ag, Al, As, Au, Be, Bi, Cd, Ce, Co, Cr, Cu, Dy, Er, Eu, Fe, Gd, Ge, Hf, Hg, Ho, In, Ir, La, Li, Lu, Mo, Nd, Ni, Pb, Pr, Pt, Ph, Ru, Sb, Sc, Sn, Sr, Ta, Tb, Te, Ti, Tl, Tm, V, W, Y, Yb, Zr,	

**Table 4.** Comparison of breaking strength and pH of soybean seeds by conventional (CF) and organic (OF) farming systems after long-term storage

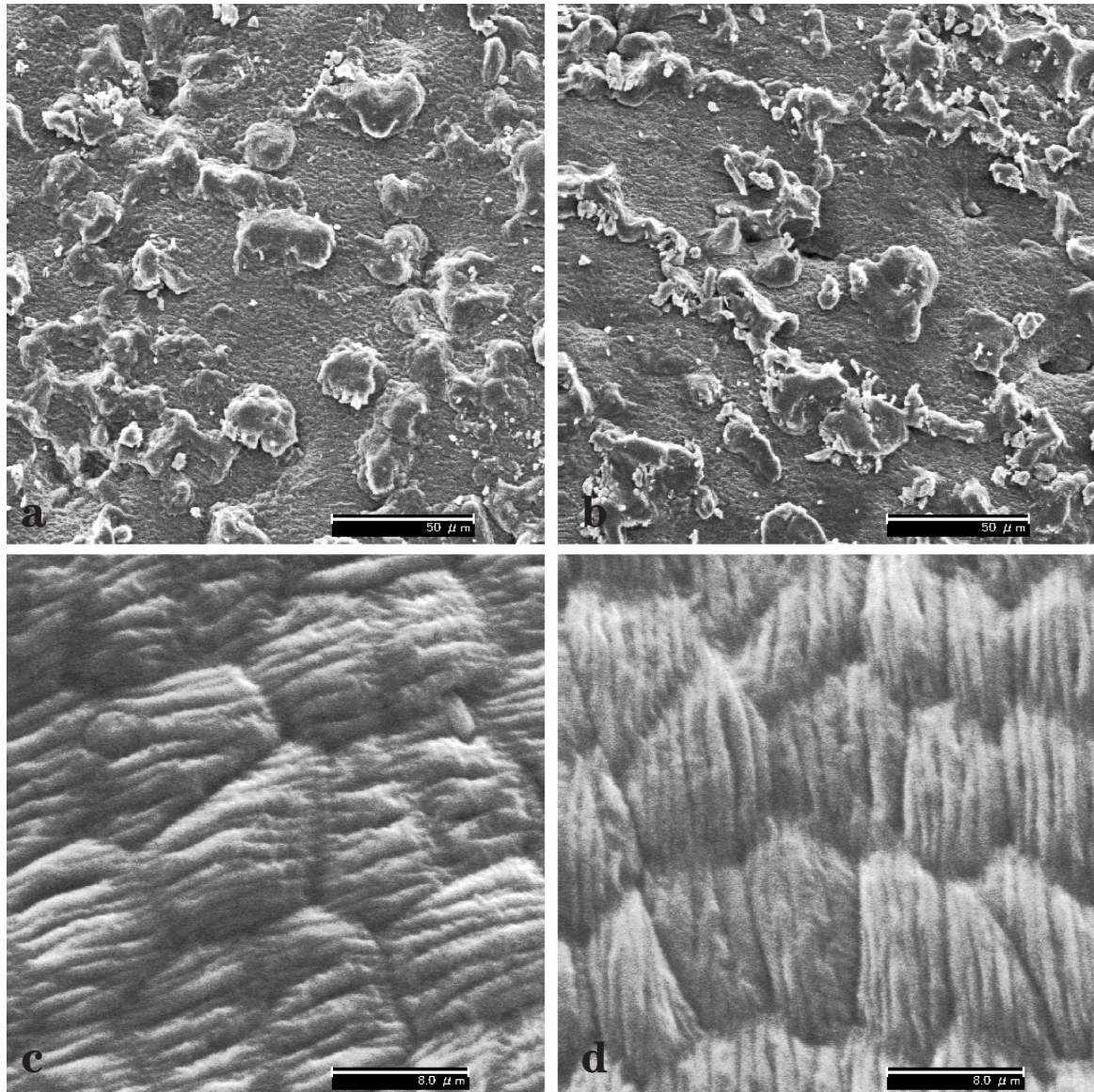
	Farming system				
	CF		OF		<i>t</i> -Test
	Mean	S.D.	Mean	S.D.	
Breaking strength (2 kgw)	16.7	3.5	17.7	2.0	$P < 0.05$
pH	6.18	0.00	6.25	0.02	$P < 0.05$

S.D.: standard deviation.

tended to be no differences in protein and lipid content of soybean seeds after harvest (**Table 1**). In addition, lipid content of soybean seeds of each group decreased about 10 % after long-term storage (**Table 1 and 2**). In the samples after long-term storage, however, the correlation ( $r = 0.8915$ ) between length and weight of CF soybean seeds was lower than that of OF seeds ( $r = 0.9211$ ) (**Fig. 3**), and the dry matter and protein content of the OF soybean seeds were higher ( $P < 0.05$  and  $0.001$ ) than CF seeds (**Table 2**). On the other hand, there were differences in the mineral compositions (more than 100 mg) of Ca, K and P content between CF and OF soybean seeds after harvest (**Table 1**). However, there was a great difference (more than 4000 mg) in mineral content, in particular K content, between CF and OF soybean seeds after long-term storage (**Table 3**).

Next, we observed the morphological cell structure

of CF and OF soybean seeds after long-term storage. By scanning electron microscopy, there was no difference in the morphological structure of the surface of the seed coat and surface of seeds which stripped off seed coat between CF and OF soybean seeds (**Fig. 4**). Cross-sections of soybean seeds were used for optical microscopy (**Fig. 1**). CF soybean seeds of soybean contained many collapsed areas in parts A–D (**Figs. 5a, c, e and g; asterisks**), whereas only few such areas were found in OF soybean seeds (**Figs. 5b and f; asterisks**). Of special note, cotyledon cells of CF soybean seeds were transformed (**Fig. 5g**). Optical microscopic analysis revealed that there were many collapsed areas and gaps in the external and internal parts of the CF soybean seeds, but not in those of OF seeds (**Fig. 5**). The storage conditions in this study were thought to be too severe (the samples were kept at ambient temperature). Nevertheless, the OF products nearly maintained their

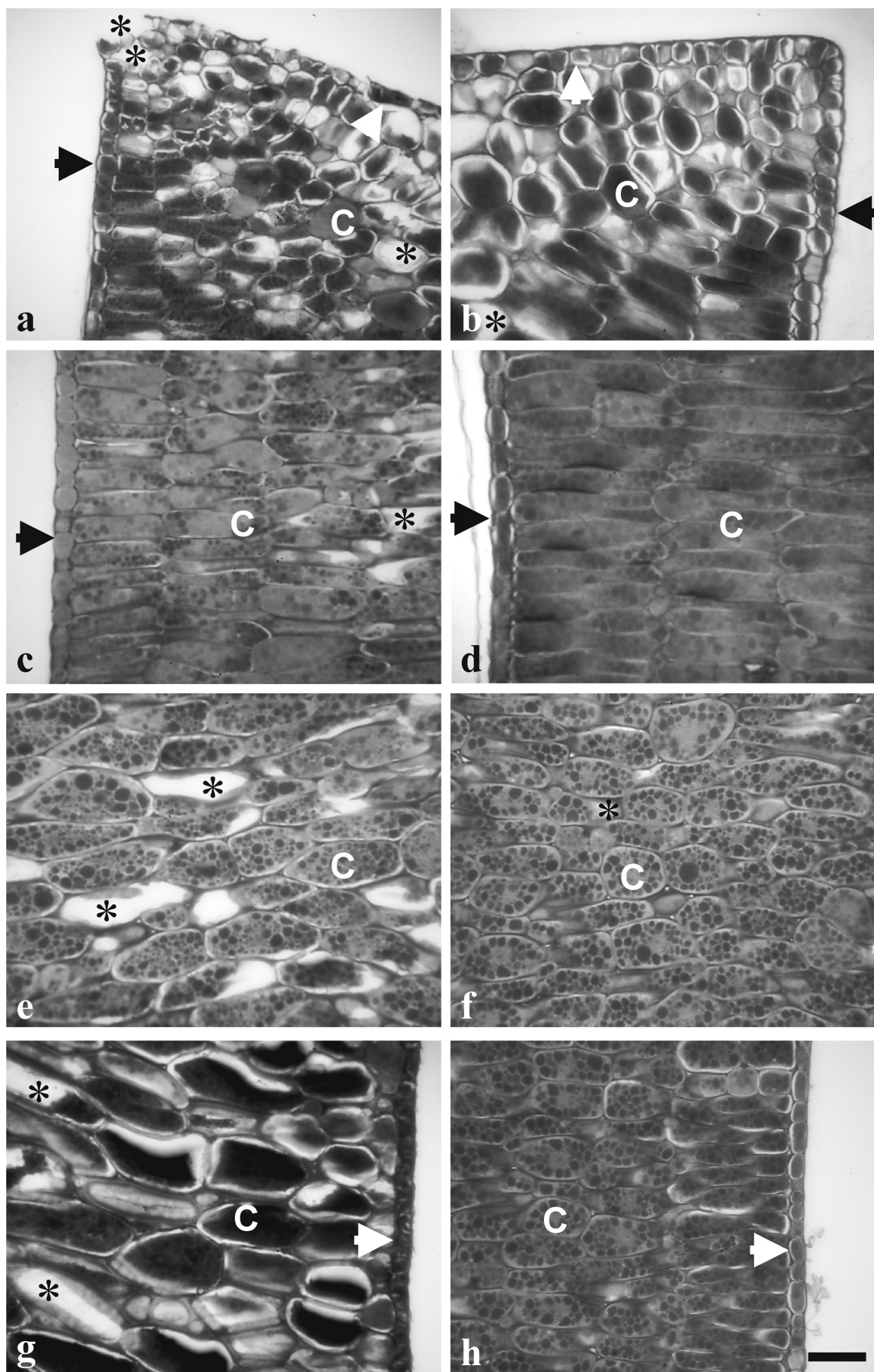


**Fig. 4.** Scanning electron microscopic photographs of surface of seed coat (a, b) and surface of seeds which stripped off seed coat (c, d) of soybean seeds after long-term (9 years) storage. a, c: CF, b, d: OF. Bar = 50 (a, b) and 8 (c, d)  $\mu\text{m}$ .

original morphology. These results indicate histological differences between CF and OF soybean seeds and suggest that CF soybean seeds became porous under storage, resulting in leakage of cell contents. Cross-sections of part C of soybean seeds were observed by transmission electron microscope (**Fig. 1**). In seeds of the CF soybean, collapsed cell walls (**Figs. 6a, e, 7c, e; asterisks**) and protein bodies (**Figs. 6c and 7a; black arrows**) were observed. The plasma membrane of the CF soybean seeds was broken, and internal organelles were observed flowing out (**Fig. 7c; white arrows**). No collapsed organelles were detected in the seeds of the OF soybeans (**Figs. 6 and 7**). Study of the OF soybean seeds revealed clear protein bodies (**Figs. 6b and 7b; PB**), lipid granules (**Fig. 7b; LG**) and thick cell walls (**Figs. 7d and f; CW**). In addition, protein bodies were systematically surrounded by clear lipid granules (**Fig. 7b**), and internal organelles were completely contained by the thick cell walls (**Fig. 7d**).

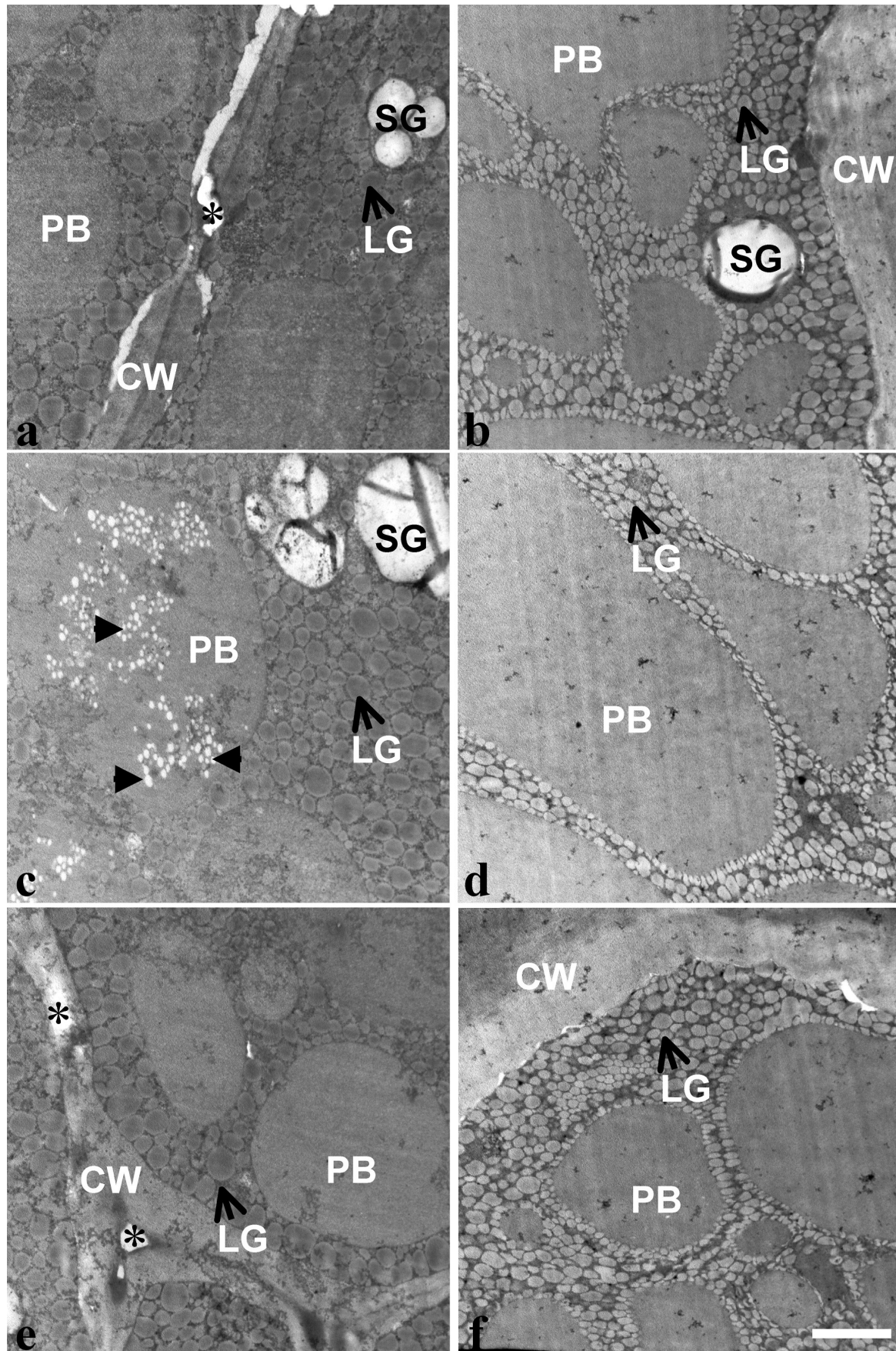
Just after harvest, Ca content of CF soybean seeds was lower (120mg) than OF ones (**Table 1**). The Ca which exists in cell wall has functions such as the maintenance of cell wall structure by combination between cell wall components (Tobias *et al.*, 1993; White and Broadley, 2003). In the cell wall of plants, Ca decreases under threshold concentration for maintaining the cell wall structure sometimes, and happen the Ca deficiency, for example cracking of the skin in tomato fruit (White and Broadley, 2003). Plants that have low Ca content show advanced ageing of tissue and increases in bacterial infections. Therefore, low molecular weight contents may escape from tissue which has low Ca content (**Fig. 5**). In this study, after 9 years of storage, the cell morphological structure (in particular, the cell wall) of CF soybean seeds collapsed (**Figs. 7c and e**). We believe that the cell wall of CF soybean seeds collapses during long-term storage because of low Ca content. The breaking strength (an index of tissue strength) of





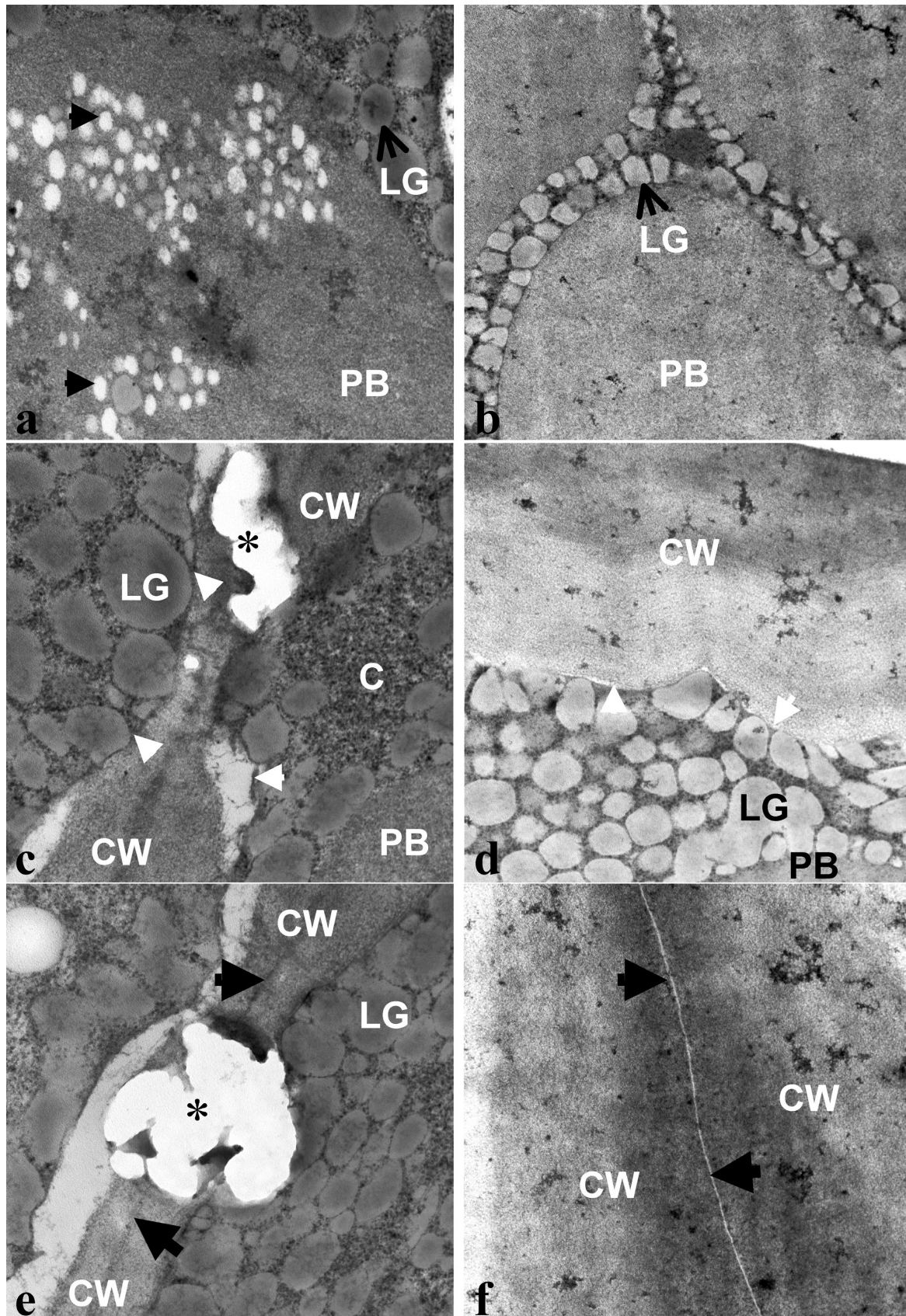
**Fig. 5.** Optical microscopic photographs of part A (a, b), B (c, d), C (e, f) and D (g, h) of soybean seeds after long-term (9 years) storage. a, c, e, g: CF, b, d, f, h: OF. C: cotyledon cells. Black arrow: inner layer, White arrow: outer layer. Asterisks mark collapsed areas in soybean seeds. Bar = 50  $\mu$ m.





**Fig. 6.** Transmission electron microscopic (TEM) photographs of part C of soybean seeds after long-term (9 years) storage. a, c, e: CF, b, d, f: OF. CW: cell wall, LG: lipid granule, PB: protein body, SG: starch granule. Asterisks: gaps and collapsed parts, Black arrows: collapsed parts. Bar =  $2\mu\text{m}$ .





**Fig. 7.** TEM photographs of part C of soybean seeds after long-term (9 years) storage. a, c, e: CF, b, d, f: OF. C: cytoplasm, CW: cell wall, LG: lipid granule, PB: protein body. Asterisk: collapsed parts, Black arrows: collapsed parts, Large black arrows: boundary between cell walls, White arrows: plasma membrane. Bar = 500 nm.



CF soybean seeds after long-term storage was lower than OF ones (**Table 4**). In addition, the loss of contents of CF soybean seeds was revealed by optical microscopy (**Fig. 5**) and a decrease of dry matter and other chemical compositions was observed (**Table 2**). In this study, there was a great difference in K contents (OF: + 4250 mg) between CF and OF soybean seeds after long-term storage in comparison with the data after harvest (**Table 1 and 3**). It is thought that the low pH of CF soybean seeds after long-term storage may be caused by the loss of K which exists mostly in the cytoplasm (**Table 4**). Conceivably, the strong cell walls of OF soybean seeds help maintain the histological structure of seeds during long-term storage. Generally, the dry weight of farming products decreases (Krishnan *et al.*, 2004), and chemical composition changes during storage (Narayan *et al.*, 1988a, b; Zhou *et al.*, 2002). However, there are no reports like this study, comparing the chemical and histological changes in CF and OF products under long-term storage. The results of this study make it clear that CF practices accelerate the deterioration of the cellular morphological properties of its products. After all, the low Ca contents of CF soybean seeds caused fragile cell walls and loss of contents in this study.

We believe that the difference in the chemical and histological characteristics between CF and OF soybean seeds was caused by the type of applied fertilizer, namely chemical or organic. It is known that the properties of soil subjected to inorganic farming differ from soil subjected to organic farming in physiochemical (Clark *et al.*, 1998; Gunapala and Scow, 1998; Liebig and Doran, 1999; Schjønning *et al.*, 2002) and biological points (Nakamura *et al.*, 2000; Fujita and Fujiyama, 2001a, b; Hirano *et al.*, 2001). The OF fields in the present study contained a rich soil fauna which in turn is likely to affect the physicochemical properties of the soil (Nakamura *et al.*, 2003). In the OF soybean field, soil properties [depth: 0 ~ 16 cm, pH: 5.35 ~ 5.66, electrical conductivity (EC): 30.9 ~ 45.5  $\mu$ S, organic carbon (Org-C): 1.8 ~ 4.8%, cation exchange capacity (CEC): 19.1 ~ 25.6 meq/100g, and base-saturation: 36.5 ~ 59.8%] differed from those of the CF fields (depth: 0 ~ 22 cm, pH: 5.60, EC: 54.7  $\mu$ S, Org-C: 1.6%, CEC: 22.0 meq/100g, and base-saturation: 36.1%). Also, the mean of the total number of enchytraeids found in soil samples from the study period (Oct. 1991 to Oct. 1998) in the OF soybean field was 549.9 individuals/m<sup>2</sup> (soil depth: 0 ~ 5 cm), in contrast with that of the CF field, which had 16.7 individuals/m<sup>2</sup> (Nakamura *et al.*, 2003). In addition, there were no earthworms in the CF field throughout the study period, but abundant earthworms were observed in the OF field (Nakamura *et al.*, 2003). These results from the present study resemble the tendencies of the soil properties of both field types (CF and OF) in other reports (Schjønning *et al.*, 2002; Lotter, 2003). Thus, both biological and physicochemical factors may affect product quality of the different farming systems (Nakamura, 1988, 1989; Woese *et al.*, 1997; Brandt and Mølgaard, 2001; Nakamura *et al.*,

2003). We guessed that the difference of change of CF and OF soybean seeds during long-term storage was caused by initial Ca contents. In another report (Kujira and Nakayama, 1997), there was a difference (OF: + 13 mg/100 g, no significance) of Ca contents between CF and OF green soybean seeds after harvest. Therefore, we consider that the absorption of Ca is inhibited by antagonism between minerals, in particular N, P, K which is present at higher levels due to chemical fertilizer in CF soybean fields.

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