



강황 수집자원의 생육특성 및 종근 저장 방법

이정훈^{1†} · 오명원² · 장현도³ · 이윤지⁴ · 정진태⁵ · 박춘근⁶

Growth Characteristics of Turmeric (*Curcuma longa* L.) Germplasms and Storage Conditions of Seed Rhizomes

Jeong Hoon Lee^{1†}, Myeong Won Oh², Hyun Do Jang³, Yun Ji Lee⁴, Jin Tae Jeong⁵ and Chun Geon Park⁶

ABSTRACT

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Background: *Curcuma longa* L., a perennial crop originating from tropical and subtropical region, including India, is noted for its important medicinal properties. However, *C. longa* plants are unable to endure the winter season in Korea, and its rhizomes were invariably succumb to fungal infection when stored in polyvinyl bags. In this study, we accordingly sought to develop a *C. longa* variety capable of producing high rhizome yields and to identify stable conditions under which rhizomes can be stored in Korea.

Methods and Results: We evaluated the agronomic characteristics of nine *C. longa* germplasms and examined the effects of storing rhizomes at different temperatures (4 °C to 24 °C) in paper bags or plastic baskets. We found that the finger rhizomes was higher in CUR02, CUR03, and CUR06 germplasms than those of other groups. Furthermore, in terms of yield per 1 m², the weights of the finger rhizomes and tuberous roots were significantly higher in CUR09 (3.4 kg/m²) and CUR04 (678.7 g/m²) than those of other groups. Therefore, we consider that these *C. longa* germplasms might be useful as breeding material. Although the fresh weights of the rhizomes were slightly reduced when stored in paper bags and a plastic baskets at 10 °C to 15 °C, there was no evidence of fungal decomposition or sprouting, which is observed when using a conventional storage method.

Conclusions: The results of this study indicate that the selected *C. longa* germplasms can provide a useful source of breeding material for the development of high yielding varieties and that a temperature ranging from 10 °C to 15 °C and the use of paper bags or plastic baskets provide stable post-harvest storage conditions for *C. longa* rhizomes.

Key Words: *Curcuma longa* L., Turmeric, Agronomic Characteristics, Storage Condition, Seed Rhizome

INTRODUCTION

Curcuma longa L. is a rhizomatous, herbaceous, perennial plant belonging to the family Zingiberaceae, and comprises approximately 70 species worldwide (Smartt and Simmonds, 1992). It is commonly distributed throughout tropical and/or subtropical regions including Cambodia, China, India, Nepal,

and etc., and particularly cultivated in India and China (Araujo and Leon, 2001). It is frequently cultivated to harvest the rhizomes (underground portion) as a medicinal herb; therefore, it is considered an important economic crop (Govindarajan and William, 1980). The deep orange-yellow powder is known as a turmeric obtained from dried its rhizomes (Li *et al.*, 2011) and used in seasoning and cosmetics. Additionally, there is a

[†]Corresponding author: (Phone) +82-43-871-5670 (E-mail) artemisia@korea.kr

¹국립원예특작과학원 약용작물과 연구사 / Researcher, Department of Herbal Crop Research, NIHHS, RDA, Eumseong 27709, Korea.

²국립원예특작과학원 약용작물과 박사 후 연구원 / Post-doc, Department of Herbal Crop Research, NIHHS, RDA, Eumseong 27709, Korea.

³국립생물자원관 연구원 / Researcher, Plant Resources Division, National Institute of Biological Resources, Incheon 22689, Korea.

⁴국립원예특작과학원 약용작물과 연구사 / Researcher, Department of Herbal Crop Research, NIHHS, RDA, Eumseong 27709, Korea.

⁵국립원예특작과학원 약용작물과 연구사 / Researcher, Department of Herbal Crop Research, NIHHS, RDA, Eumseong 27709, Korea.

⁶국립원예특작과학원 약용작물과 연구관 / Researcher, Department of Herbal Crop Research, NIHHS, RDA, Eumseong 27709, Korea.

specific interest in turmeric because of its reported medicinal potential (Cousins *et al.*, 2007).

A total of 235 compounds have been detected in the leaves, flowers, roots, and rhizomes of *C. longa*, including 22 diarylheptanoids and diarylpentanoids, 8 phenylpropene and other phenolic compounds, 68 monoterpenes, 109 sesquiterpenes, 5 diterpenes, 3 triterpenoids, 4 sterols, 2 alkaloids, and 14 other compounds (Li *et al.*, 2011). Turmeric also contains curcumin, which confers a yellow color (Jayaprakasha *et al.*, 2002). Turmeric oil is reported to possess biological activity, including carminative, anti-fungal, and anti-platelet effects (Lee, 2006). Additional reports attribute anti-inflammatory, anti-cancer, anti-tumor, anti-viral, and hepatoprotective activity to turmeric (Ammon and Wahl, 1991). In addition to the above-mentioned activities, *C. longa* has pharmacological activities such as antioxidant, anticoagulant, anti-diabetic, anti-microbial, anti-ulcer, wound healing, and anti-fertility (Yadav and Tarun, 2017).

The underground part of *C. longa* is divided into three parts: Finger rhizome, tuberous root, and root. The *C. longa* rhizome which is composed of a mother rhizome with primary, secondary, and tertiary fingers, is very large, yellow when horizontally cut, and has a unique scent (An, 2000). *C. longa* typically demonstrates a nutritional growth period of 7 to 8 months and undergoes a flowering period for 2 to 3 months. The rhizome is formed when flowering begins and it enters a period of dormancy when flowering stops and the aerial portion withers.

The humid climate and temperature between 20°C and 30°C are needed for *C. longa* cultivation. It grows well under rich sunshine in fertile, sandy soil, that is well drained. However, *Curcuma* spp. plants including *C. longa* do not grow naturally in Korea. *C. longa* is produced as an annual plant by using rhizomes because a “seed” in Korea as seeds are not produced in its life cycle.

When *C. longa* are cultivated in Korea, the resulting rhizomes are considered inappropriate for medicinal use due to low pharmacological composition of curcumin. In addition, Korean environmental conditions also result in a reduced yield of rhizome. Therefore, to produce a high yield of rhizome with improved quality, further studies of *C. longa* development are needed.

The rhizome does not survive well in winter owing to cold injury resulting from the low temperatures in Korea (Lim *et al.*, 2013), and thus it is important to consider the optimal storage methods employed during the winter season. One study

suggested that planting needs to occur at the end of April in southern parts in Korea, due to the thermophilic properties of this crop (Choi, 2004). Rhizomes were stored by packing in polyvinyl bags in a warm area or on the ground and sprayed with water at intervals of a few days. However, this method was not proven and resulted in the growth of fungi. Another report suggested packing *C. longa* rhizomes with vermiculite and maintaining the temperature above 10°C (Lim *et al.*, 2013). Vermiculite absorbs moisture initially but its potential to cause corruption is a concern.

In this study, we evaluated several *C. longa* genetic resources to determine their agronomic traits and to utilize as a breeding material. Additionally, we evaluated different storage temperatures and containers in an attempt to discover stable rhizome storage methods which do not involve curing.

MATERIALS AND METHODS

1. Plant materials

The *Curcuma longa* L. genetic germplasms used for evaluation of growth characteristics comprised a total of 9 germplasms (Table 1). The materials were collected in Korea, China, and Japan and was provided from the National Agrobiodiversity Center (Jeonju, Korea). These were compared with *C. longa* cultivated in Jindo of Jeollanamdo as a control, which is known as local species (Jindo henceforth). In addition, *C. longa* seed rhizomes were divided into three or four for evaluation of different storage conditions as outlined below.

The test field formed ridges and covered with black plastic bags after the compost was applied using basal fertilization at a total of 4,000 kg per 10 a, considering the soil fertility. The experiment plot was arranged by the randomized complete block design with triplicate. For planting density, furrow spacing was 30 cm in the 90 cm ridges, with 20 cm of plant intervals. The yield was converted to the yield per 10 a after harvesting in 1 m² experiment plot.

2. Agronomic characteristics

C. longa germplasms were cultivated in a National Institute for Horticulture and Herbal Science (NIHHS) experimental field located in Eumseong according to the standard cultural practices. The emergence rate was investigated after 50 days.

The harvested *C. longa* plants were examined for agronomic characteristics including plant height, leaf length, leaf width,

Table 1. *Curcuma longa* germplasm used in this study.

Accession No.	Collected country
<i>Jindo</i> ¹⁾	Korea
CUR01	China
CUR02	Japan
CUR03	Japan
CUR04	Japan
CUR05	Japan
CUR06	Japan
CUR07	Japan
CUR08	China
CUR09	China

¹⁾*Jindo*; local species cultivated in Jindo used as control in this study.

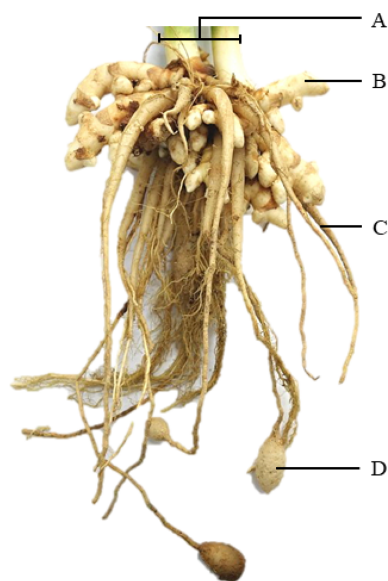


Fig. 1. Characters of the underground portion of *C. longa*. A; stem width, B; finger rhizome, C; root, D; tuberous root.

finger rhizome weight, tuberous root weight, etc. The underground portion was also divided into 2 sections and examined (Fig. 1). Investigations were performed from 2015 to 2016 and three independent experiments were performed, representing biological replicates. All data had shown the average of 2 years.

3. Storage conditions

Rhizomes were placed in a paper bag and a plastic basket, respectively. And they were stored at 4°C, 10°C, 15°C, and 24°C (room temperature, RT hereafter) for 30, 60, and 90 days.

The conventional storage method (in sealed polyvinyl bags

at 4°C, sprayed with water every 2 days) was compared as a control. Every 30 days, the loss of fresh weight, the level of fungal contamination, and sprouting rate were recorded. This experiment had a randomized complete block design. Three independent experiments were performed, representing biological replicates.

4. Statistical analysis

Statistical significance of the results was evaluated using One-way ANOVA for comparisons between multiple groups. All calculations were performed using SPSS software Version 22.0 (IBM Co., Armonk, NY, USA). Differences were considered statistically significant at a *p* value of < 0.05.

RESULTS AND DISCUSSION

1. Agronomic characteristics of germplasms

To develop new *Curcuma longa* L. varieties with improved yield and quality and adapted to a Korean cultivation environment, growth characteristics including emergence rate for aerial and underground portions of *C. longa* plants were evaluated.

Fig. 2 showed the appearance of the underground portion of *C. longa*. Emergence rate was examined from 50 days after rhizome transplantation. The date of sprouting was earlier for CUR02 and CUR03 than for the control *Jindo*, however, the overall emergence rate was above 95%, indicating that these resources had the high germination rate (Table 2).

There was no significantly difference among 9 germplasms during the initial growth stage, while during the later growth stage, CUR04 had the longest plant height. The weight of the aerial portion was 584.1, 577.5, and 610.3 g/plant in CUR03, CUR04, and CUR06, respectively, which were higher than that of other germplasms (Table 3).

Yield per unit area was determined by calculating the number of plants and the weight of root, finger rhizome, and tuberous root. The weight of finger rhizome among individuals in CUR02, CUR03, and CUR06 was 222.9, 222.5, and 225.9 g/plant, respectively, which were higher than those of other groups. The tuberous root weight was the highest in CUR04, which was 23.5 g (Table 4).

On the other hand, when compared to the yield per 1 m², a yield of finger rhizome was significantly higher in CUR09 (3.4 kg/m²), CUR01 (3.3 kg/m²), and CUR06 (3.3 kg/m²), and that of tuberous root was significantly higher at CUR04 (678.7 g/m²)

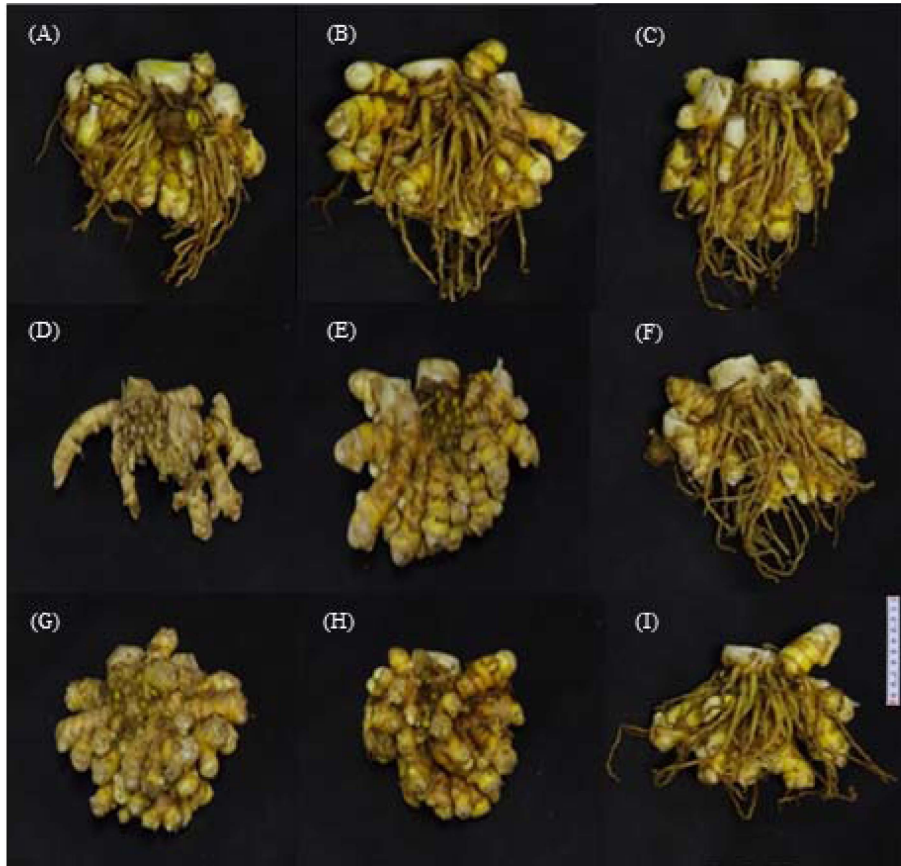


Fig. 2. Characteristics of the underground portion of *C. longa* germplasm. (A); CUR01, (B); CUR02, (C); CUR03, (D); CUR04, (E); CUR05, (F); CUR06, (G); CUR07, (H); CUR08, (I); CUR09.

Table 2. Emergence rate (%) of germplasm.

Line	<i>Jindo</i> ¹⁾	CUR01	CUR02	CUR03	CUR04	CUR05	CUR06	CUR07	CUR08	CUR09
50 days after planting (June 17)	12	5	22	20	18	16	7	12	5	5
60 days after planting (June 26)	61	39	72	64	57	59	44	50	54	38
70 days after planting (July 07)	76	65	82	82	75	74	61	71	71	63
Average of emergence rate (%)	99.3 ^a	98.6 ^a	97.2 ^a	97.8 ^a	98.6 ^a	98.0 ^a	96.5 ^{ab}	95.9 ^{ab}	98.6 ^a	97.8 ^a

¹⁾*Jindo*; local species cultivated in Jindo used as control in this study. *Alphabet means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

among 9 germplasms (Table 5). However, there was no significant correlation between finger rhizome and tuberous root, it is considered that the yield was decided by individual characteristics or environmental difference.

Assessing genetic resources is considered to be crucial for the genetic improvement of crops. Evaluation of genetic diversity assists in improving the understanding of the genetic background and in demonstrating the breeding value of different germplasm lines.

Plant height and number of leaves have been reported to be

determinants of yield potential in individual genotypes in the context of improving turmeric crops (Narayanpur and hanamashetti, 2003).

Therefore our study might be used as fundamental data for breeding variety through the evaluation of the agronomical characteristics of collected *C. longa*.

2. Storage conditions of seed rhizomes

This study was conducted to determine the optimal conditions for storage of rhizomes during the winter season.

Table 3. Growth characteristics on aerial portion of *C. longa* germplasm.

Line	Investigating time	Plant height (cm)	Leaf			Stem		Aerial portion weight (g/plant)
			Number (ea/plant)	Length (cm)	Width (cm)	Number (ea/plant)	Width (mm)	
Jindo ¹⁾	early in Aug.	59.7	4.9	26.8	12.4	0.4	14.5	410.1±123.6 ^b
	the end of the Oct.	132.5	7.7	56.7	17.6	1.9	31.6	
CUR01	early in Aug.	54.5	4.4	25.7	13.4	0.1	13.8	389.7±41.6 ^b
	the end of the Oct.	137.4	7.7	54.1	18.4	1.8	32.6	
CUR02	early in Aug.	57.8	5.0	25.8	12.1	0.2	15.4	517.7±96.7 ^{ab}
	the end of the Oct.	137.8	7.7	52.0	17.1	1.7	34.8	
CUR03	early in Aug.	60.8	5.1	27.9	12.3	0.4	15.5	584.1±199.3 ^a
	the end of the Oct.	137.2	7.5	53.0	17.4	1.4	33.8	
CUR04	early in Aug.	57.6	5.4	25.8	12.0	0.7	14.0	577.5±55.3 ^a
	the end of the Oct.	143.2	8.1	53.7	17.1	1.8	27.2	
CUR05	early in Aug.	58.3	5.0	26.7	12.2	0.2	14.8	482.3±55.3 ^{ab}
	the end of the Oct.	136.0	7.6	54.6	17.6	1.7	32.8	
CUR06	early in Aug.	56.9	4.8	26.0	11.8	0.1	14.3	610.3±235.0 ^d
	the end of the Oct.	136.7	7.7	56.8	18.0	2.0	34.4	
CUR07	early in Aug.	53.8	4.8	25.3	11.5	0.2	13.7	421.2±62.8 ^b
	the end of the Oct.	132.0	7.6	52.0	16.8	1.8	33.3	
CUR08	early in Aug.	54.5	4.7	25.1	11.3	0.1	14.4	418.4±28.9 ^b
	the end of the Oct.	131.1	7.7	55.4	17.3	1.6	31.3	
CUR09	early in Aug.	52.1	4.7	24.7	11.2	0.1	13.5	407.9±78.0 ^b
	the end of the Oct.	128.5	7.6	50.4	16.5	1.7	32.1	

¹⁾Jindo; local species cultivated in Jindo used as control in this study. *Alphabet means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

Table 4. Growth characteristics on underground portion of *C. longa* germplasm.

Line	Finger rhizome			Tuberous root			Root		
	Length (cm/ea)	Width (mm/ea)	Weight (g/plant)	Length (cm/ea)	Width (mm/ea)	Weight (g/plant)	Length (cm/ea)	Width (mm/ea)	Weight (g/plant)
Jindo ¹⁾	7.1±0.5	22.3±0.5	211.8±57.0 ^a	5.9		15.6 ^{ab}	15.0±2.1	6.3±0.2	36.4±12.3 ^b
CUR01	7.7±0.6	22.2±0.5	198.2±43.9 ^a	3.2		9.1 ^{ab}	14.5±2.2	6.6±0.5	42.6±13.5 ^b
CUR02	7.5±0.4	23.3±0.1	222.9±51.1 ^a	3.9±0.1	13.7±0.9	11.2±4.2 ^{ab}	14.2±2.1	7.2±0.1	47.5±13.9 ^b
CUR03	7.6±0.4	22.8±0.2	222.5±23.4 ^a	3.1±0.1	8.6±1.3	8.0±6.0 ^{ab}	14.8±1.4	6.6±0.2	43.8±12.4 ^b
CUR04	9.3±0.3	20.8±0.4	163.2±37.1 ^a	3.9±0.7	9.7±5.4	23.5±20.8 ^a	18.5±0.0	8.9±0.3	74.7±9.5 ^a
CUR05	7.5±0.3	22.8±1.0	208.6±48.8 ^a	4.0±0.0	11.7±0.6	9.1±4.1 ^{ab}	14.6±1.6	6.6±0.4	38.8±9.9 ^b
CUR06	7.5±0.6	23.3±1.0	225.9±40.0 ^a	4.4±0.7	14.7±1.4	11.0 ^{ab}	14.4±0.6	6.3±0.4	37.0±6.3 ^b
CUR07	7.3±0.3	22.0±0.1	171.5±41.4 ^a	4.3±0.1	12.2±0.5	10.1±6.4 ^{ab}	15.7±0.4	6.5±0.2	37.2±14.4 ^b
CUR08	7.2±0.4	22.3±0.1	187.7±50.9 ^a	4.0	11.9	7.8±2.3 ^{ab}	14.5±0.6	6.5±0.3	35.2±7.8 ^b
CUR09	7.2±0.6	21.6±0.8	201.8±61.2 ^a	4.8	14.8	5.3±1.8 ^b	14.2±1.5	6.6±0.3	36.7±10.1 ^b

¹⁾Jindo; local species cultivated in Jindo used as control in this study. *Alphabet means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

Table 5. Yield traits per 1 m² on finger rhizome and tuberous root of *C. longa* germplasm.

Line	Finger rhizome weight (kg/m ²)	Tuberous root weight (g/m ²)
<i>Jindo</i> ¹⁾	3.1±0.1 ^{ab}	213.3±50.5 ^b
CUR01	3.3±0.2 ^{ab}	177.0±24.0 ^b
CUR02	3.2±0.5 ^{ab}	251.0±40.2 ^b
CUR03	3.0±0.2 ^{ab}	166.0±68.2 ^b
CUR04	2.3±0.3 ^b	678.7±131.5 ^a
CUR05	3.2±0.2 ^{ab}	242.3±35.2 ^b
CUR06	3.3±0.4 ^{ab}	192.3±27.3 ^b
CUR07	2.7±0.3 ^{ab}	206.3±24.5 ^b
CUR08	3.2±0.1 ^{ab}	159.0±48.2 ^b
CUR09	3.4±0.3 ^a	131.7±32.1 ^b

¹⁾*Jindo*; local species cultivated in Jindo used as control in this study. *Alphabet means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

Fresh weight loss, decomposition rate attributed to fungi and sprout rate were evaluated according to the storage containers and temperature.

When rhizomes were stored in a polyvinyl bag at 4°C, as in a conventional method, fresh weights increased due to water absorption. However, when stored in a paper bag and basket container, there was no significant change. At 10°C and 15°C, there was no remarkable difference or a slight decrease whereas at RT conditions (24°C), fresh weight markedly decreased when stored in paper bags and basket containers (Fig. 3). These results suggest that paper bags and basket containers are better than polyvinyl bags for rhizome storage.

When *C. longa* rhizomes were stored at 4°C in a polyvinyl bag, fungal decomposition rate was significantly higher, suggesting that fungal growth occurred due to the addition of water (Fig. 4). When ginger rhizomes were stored in polyethylene file, the weight loss ratio was lower than that of the control (untreated) rhizomes; however, decomposition rate and incidence of fungi were higher (Choi and Kim, 2001). Bambirra *et al.* (2002) also reported that polyethylene bags were not suitable material for storage of the ground turmeric due to allowing absorption of moisture. This result was shown the same tendency to our present study, which rhizomes stored in polyvinyl bag were absorbed moisture and it induced the fungal decomposition.

Among various physiological factors affected by storage,

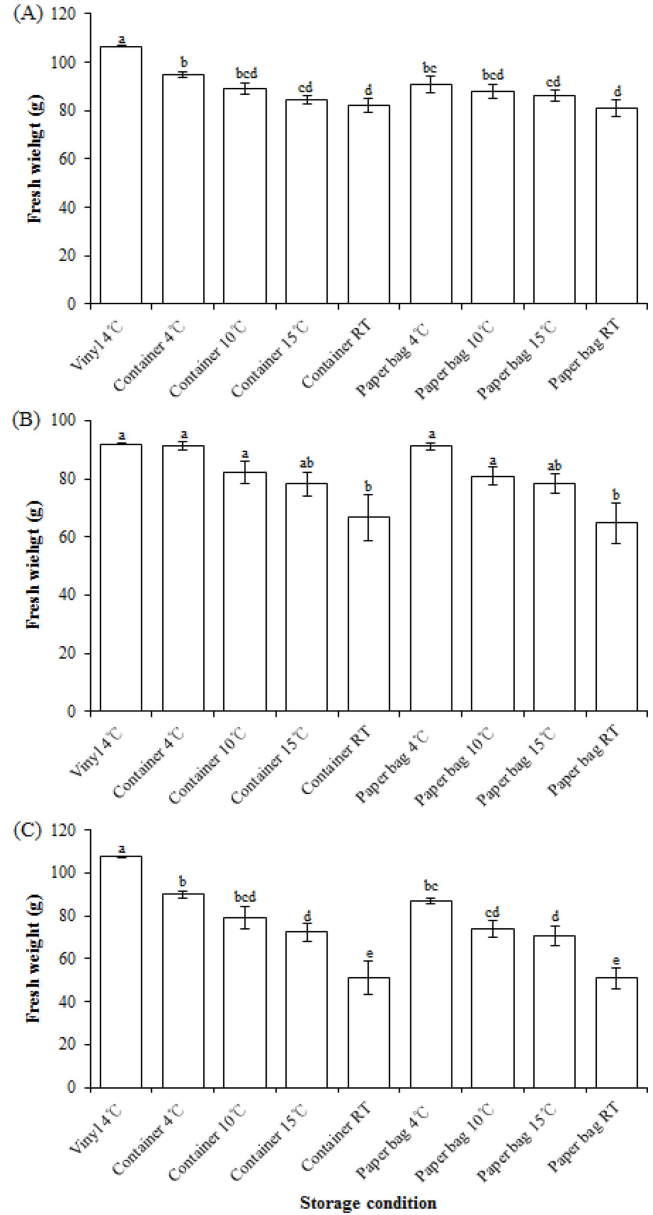


Fig. 3. Changes in fresh weight (g) of *C. longa* rhizome exposed to different storage conditions. The storage duration of *C. longa* rhizome was 30 days, 60 days, and 90 days, respectively, which shown (A), (B), and (C). Data are shown as means ± SD from three independent biological replicates. *Means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

sprouting is considered the most noticeable symptom of deterioration in the case of many tuber crops (Thomas, 1988; Jaleel *et al.*, 2007). *C. longa* rhizome sprouting was observed during storage at RT conditions (Fig. 5), induced by the higher temperature. These results indicate that 4°C and RT conditions

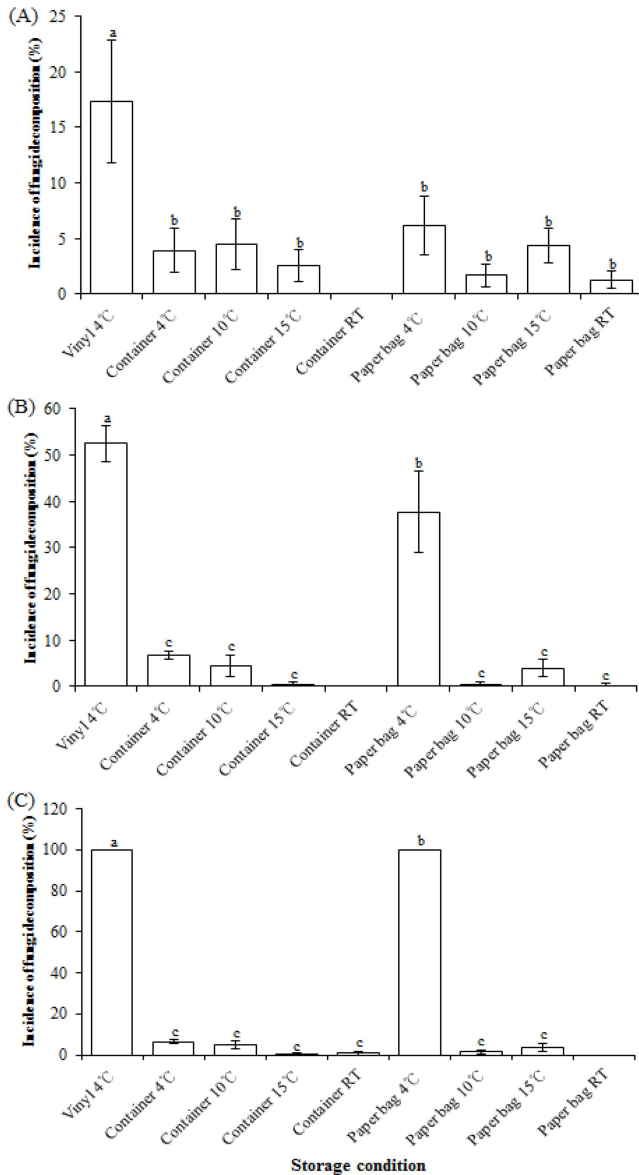


Fig. 4. Incidence of fungal decomposition in *C. longa* rhizome exposed to different storage conditions. The storage duration of *C. longa* rhizome was 30 days, 60 days, and 90 days, respectively, which shown (A), (B), and (C). Data are shown as means \pm SD from three independent biological replicates. *Means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

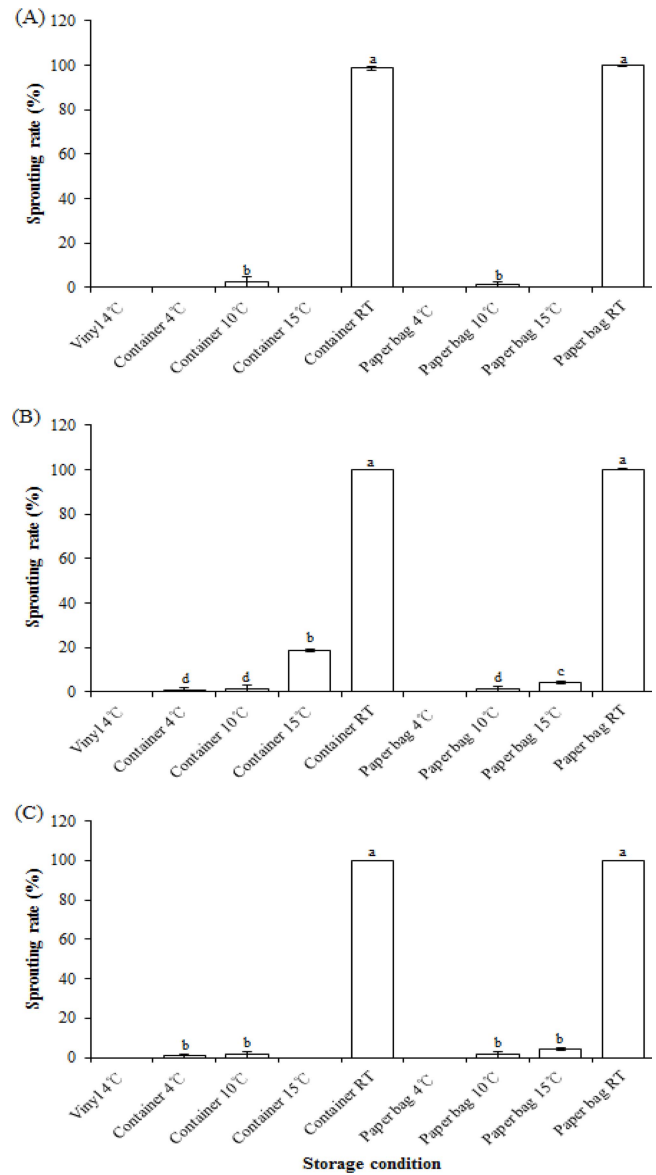


Fig. 5. Sprouting rate in *C. longa* rhizomes exposed to different storage conditions. The storage duration of *C. longa* rhizome was 30 days, 60 days, and 90 days, respectively, which shown (A), (B), and (C). Data are shown as means \pm SD from three independent biological replicates. *Means with the same letter are not significantly different according to Duncan's Multiple Range Test (DMRT, $p < 0.05$).

are not suitable for rhizome storage due to risk of fungal decomposition (Fig. 6) and sprout emergence, respectively. Thus, these results suggest that *C. longa* rhizomes remain stable when stored at a temperature between 10°C and 15°C in a paper bag and basket container.

Similarly, tissue softening, root discoloration, and decay were observed in ginger rhizomes stored below 12°C (Akamine,

1962). Lim *et al.* (2013) reported that the optimal storage temperature for *C. longa* rhizomes was above 10°C. This is similar to the results of our study. Taken together, these results suggest that temperatures between 10°C and 15°C provide suitable storage conditions for rhizomes without resulting in chilling-damage.

Harvested roots, tuber, and bulb crops demonstrated

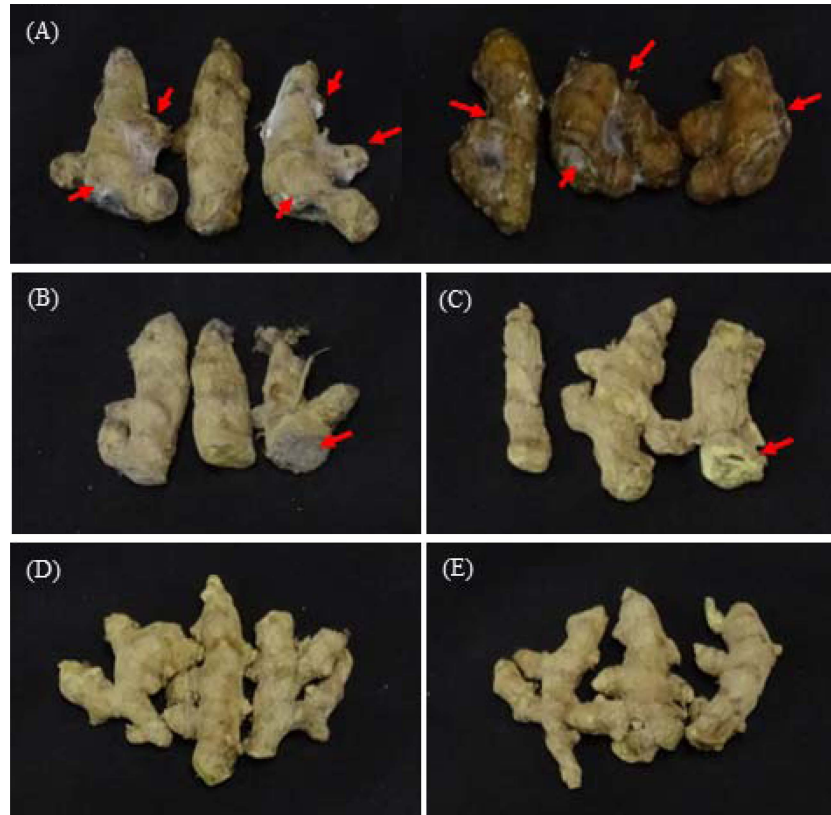


Fig. 6. The contamination of fungi after storage of *C. longa* rhizome with different storage temperatures and containers at 90 days. Arrows means the contamination of fungi after storage. *C. longa* rhizome was stored at 4°C in polyvinyl bag as a practice method (A), 10°C in paper bag (B), 10°C in plastic basket (C), 15°C in paper bag (D), 15°C in plastic basket (E), respectively.

decreased weight and decomposition during the storage period; therefore, a simple and effective curing process is required for these plant parts (Cantwell and Kasmire, 1992). Curing protects the rhizomes from contamination at the cutting stage, and controls respiration, moisture transpiration, and sprout development (Salunkhe and Desai, 1984). However, sometimes curing introduces more fungi associated with cutting injury from the harvesting process (Kim *et al.*, 2012). In a study of weight loss and sprouting control in ginger rhizomes, waxing was not shown to reduce moisture content (Paull *et al.*, 1988). Therefore, *C. longa* rhizomes need to be stored without curing similar to ginger.

In this study, 9 lines of *C. longa* were evaluated to provide breeding materials for suitable variety in Korea. Based on growth characteristics, among 9 germplasms, the weight of finger rhizome on CUR02, CUR03, and CUR06 were higher than that of others. In addition, CUR09 was shown the high tendency of finger rhizome on yield per 1 m², on the other hand, CUR04 was shown the significantly highest yield per 1

m² of tuberous root. Therefore, it suggest that these germplasm such as CUR09 and CUR04 might be selected useful germplasm for breeding material of finger rhizome and tuberous root, respectively. Furthermore, when *C. longa* rhizome was stored after harvest, there was some problem. To establishment of optimal storage condition, rhizomes were placed to paper bags, plastic baskets, and polyvinyl bag as a control under different condition. There was not observed any evidence of fungal decomposition or sprouting in rhizome which stored in paper bags or plastic basket at 10°C and 15°C compared to a polyvinyl bag at 4°C.

These results suggest that the *C. longa* rhizome will be able to be stably stored under storage temperature of 10°C to 15°C in paper bags or plastic baskets.

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