

**MYCOFLORA OF GRAIN MAIZE (*Zea mays* L.) STORED IN
TRADITIONAL STORAGE CONTAINERS (GOMBISA AND SACKS) IN
SELECTED WOREDAS OF JIMMA ZONE, ETHIOPIA**

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ABSTRACT

Mycoflora of maize (*Zea mays* L.) grain (Variety: Bako Hybrid-660) stored in two traditional storage containers (*Gombisa* and Sacks) for 180 days was studied for mycoflora in two agro-ecologies, that is Intermediate and Lowland, with altitude ranges of 1500-2500 meters above sea level, and 1000-1500 meters above sea level, respectively, in Jimma zone, Ethiopia. The temperature and relative humidity were observed for identifying fungi species which can flourish and cause maximum deterioration to maize grains. Significant ($P < 0.05$) decreases in germination rate of the grains were observed with time under each storage method for both the low and intermediate altitude ranges. Germination percentage reduced from 98% and 97.5% to 68.5% and 80.5% for grains stored in *Gombisa* and Sacks, respectively. Storage type significantly ($p < 0.05$) affected seed germination under intermediate agro-ecology whereas no significant ($p > 0.05$) effect was observed under lowland agro-ecology due to storage container type. One sterile white mycelium and a total number of eight species of fungi viz., *Aspergillus flavus*, *Aspergillus fumigatus*, *Aspergillus niger*, *Aspergillus terreus*, *Cladosporium cladosporioides*, *Drechslera halodes*, *Fusarium oxysporum* and *Penicillium chrysogenum* were identified from maize grain at the beginning and during storage. The most common fungi recorded from both agro-ecologies in the two storage structures were *A. flavus*, *A. niger*, *D. halodes* and *F. oxysporum*. These fungi were recorded from 90, 51, 72 and 44 percentage of the seed samples, respectively. In the districts of both intermediate and lowland agro-ecology these fungi were consistently recorded throughout 180 days, during storage. *A. fumigatus* was detected in 3.6% of samples while *C. cladosporioides* observed in 15% of the samples. Fungal species *A. terreus* and *Penicillium* were recorded in 0.5% of samples in selected districts of Jimma. These fungal species were known to cause deterioration of maize and are a health risk to humans and animals due to the toxins they potentially produce.

Key words: Agro ecologies, Fungi, Maize, Storage

INTRODUCTION

In Ethiopia, maize (*Zea mays* L.) is a staple food and one of the main sources of calories in the major producing areas [1]. The crop ranks first in total production area and yield per hectare [2]. Maize has been selected as one of the national commodity crops to satisfy the food self-sufficiency program of the country to feed the alarmingly increasing population [1]. However, the grain undergoes quantitative and qualitative losses during storage. The losses occur mainly because of improper storage [3]. A large number of pathogenic fungi, bacteria, viruses and insects infecting and infesting maize grain cause combined worldwide annual losses of 9.4% [4]. Fungi affect the quality of grain as a result there will be, increase in fatty acid, reduction in germination, increase its mustiness, production of toxins and finally leading to spoilage of grain in many ways.

Fungi are the second important cause of deterioration and loss of maize next to insects [5, 6]. Fungi could cause about 50 to 80% of damage on farmers' maize during storage if conditions are favorable for their development [5]. The major genera commonly encountered on maize in tropical regions are *Aspergillus*, *Fusarium* and *Penicillium* [7, 8]. This is the cause of concern because these genera have species capable of producing a wide spectrum of compounds shown to be toxic to man and animals [9 – 15].

A survey conducted in three major maize grain producing areas of Ethiopia indicated that the majority of farmers (93.3%) use traditional storage containers that expose their stored grains to attack by storage pests and/or other factors. The average actual loss per household was about 12 percent of the average total grain produce [16]. Since deterioration of stored grains results from the interactions of pathogens, physical factors and storage containers it is important to understand the inter-relations and interactions of these variables in order to design effective control and management of these factors for safe grain storage [17]. Grain storage containers being used by majority of farmers in Jimma zone (more than 97%) are traditional ones that couldn't protect the stored grain from deterioration [18]. There is no information on the exact cause of deterioration of grains stored in these traditional storages in the zone that could serve as basis to take corrective measures. Availability of such information will help to take corrective measures in improving post harvest grain management and hence support the efforts being made to become a food secured country. The patterns of storage temperature, relative humidity and the associated fungi of maize stored in the traditional storage containers in the area are not clearly known and documented. It is, therefore, essential to study the existing fungal species that infect grains of maize stored in two major traditional storage containers (*Gombisa* and Sacks) and the influence of altitude/elevation (both intermediate and lowland agro-ecologies) on fungal infection of stored maize grain over time under traditional storage facilities.

MATERIALS AND METHODS

Description of the study area

The study was carried out in Jimma zone, Ethiopia which is found at about 345 km from Addis Ababa in South west and lies between 36° 10' E longitude and 7° 40' N latitude. The zone has an elevation ranging from 880 to 3360 meters above sea level (masl). The area experiences annual average rainfall of 1000 mm for 8 to 10 months. The main rainy season extends from May to September and the small rainy season takes place in February, March and April. The temperature of Jimma zone varies from 8-28°C. The average annual temperature is 20°C [19].

The agro-ecologies of the study area have an altitude range of 1000-1500 (lowlands), 1500-2500 (intermediate) and 2500-3360 masl (highlands) [20]. Only two agro-ecologies (intermediate and lowlands) growing BH-660 maize variety were selected for the study since BH-660 maize variety is currently not produced in the highland agro-ecology of the study area. Districts that exhibit wide production of BH-660 maize variety in the study area were selected, Omo Nada and Kersa from intermediate agro-ecology and Tiro Afeta and Sekoru from the lowland agro-ecology.

Experimental design and treatments

Factorial arrangement using Completely Randomized Design (CRD) was employed for the experiment in two replications. There were three factors: traditional maize storage containers at two levels (*Gombisa* and Sack), agro-ecologies at two levels (lowland and intermediate), and storage periods at four levels (initial, 60, 120 and 180 days after harvest). The study was conducted for six months in 2009/2010 harvesting season by taking samples at two months interval. Baseline data was collected at the start of the study period for comparison. A total of 96 samples (12 samples per treatment) were taken during six months of the storage period.

Experimental materials

The experimental materials used for the study were BH-660 variety of maize grain harvested in December 2009 and two types of traditional maize storage containers (*Gombisa* and Sacks).

Gombisa is an above ground cylindrical shaped traditional storage bin used to store maize cob. It is vertical un-plastered wall made from woven plant stems usually Eucalyptus. Conical grass- thatched roofing is used to cover the top end, usually overhanging it about 70 cm out of the wall. The structure rests on a low level (10 to 60 cm) wooden platform mounted on four, six or eight stone pillars or on a short yoked wooden post (10 to 60 cm) firmly driven in to the ground (penetrating the ground). The floor is made of mats of woven bamboo splits and dry stalks of sorghum under which a wooden beam of Eucalyptus laid. The floor is usually plastered with salt and cow dung mixture (1.5 kg table salt per kg cow dung and a liter of water). The height of *Gombisa* ranged from 124 to 155 cm while its diameter varies from 148 to 304 cm. consequently the capacity varies from 10 to 25 Quintals of maize cobs. It

is generally neither air tight nor moisture and insect proof. Maize cob loading and unloading are effected by removing the thatched roofing.

A polypropylene Sack is the other type of storage container to store shelled maize after treating it with either of traditional or modern pesticides. It is made of woven synthetic fiber and low cost indoor storage container that can hold 50 to 100 kg of shelled maize grain which is a readily available. Rodents are known to dislike nibbling polypropylene Sacks, as the strands tend to get in between their teeth (21). It cannot be used for storing maize for long term storage (more than 6 months) because the material tends to inhibit free circulation of air with in the Sack. Polypropylene Sack is also difficult to fumigate. It is puncture resistant and has moderate permeability to moisture, gases and odors (21).

Sampling of the Grain for Evaluation

Cob samples in *Gombisa* were placed in three meshed wire cages that can hold about 10 cobs each. The cages were placed at the bottom end of 11 cm diameter polyvinyl chloride (PVC) tube the surface of which is perforated with 1.4 cm diameter so that the environment in the cage and the PVC tube is identical to that in the rest of the *Gombisa*. The cobs placed in the cages were each attached to a string the other end of which appears at the top of the stored grain. This arrangement is to facilitate removal of the sample cobs for analysis during the storage period. Three such cages connected to PVC tubes were placed at the center and the two sides of the middle layer in each *Gombisa* filled with maize cobs. They serve as representative samples of the stored maize cobs for evaluation throughout the study period.

Initial samples of six cobs were randomly taken before the bulk was loaded in to *Gombisa*, shelled manually to make 1 kg and then kept in an air-tight plastic bag. The initial maize samples from each storage containers were taken as a control at the beginning of the storage period. Subsequent sampling from *Gombisa* and Sacks was carried out at an interval of two months for the storage period of six months. Three cobs were drawn from each cage via the tube using strings, shelled manually and thoroughly mixed. For sampling grain from the Sacks, the procedure described in AOAC (1995) was followed [22].

Physical Variables

Grain moisture content

The grain moisture content was measured using a calibrated moisture tester (Dickey-john Corp. Auburn, IL 62615 USA) immediately before storage and during the subsequent grain sampling periods from each storage container.

Storage temperature and relative humidity

The storage temperature and relative humidity were measured at an interval of 15 days during the storage periods using portable digital Thermo- Hygrometer (Hanna, HI8564, Italy) [23].

Seed germination test

Seed germination test was conducted using standard procedures of ISTA (1996) [24].

Identification of fungi

A total of sixty maize seeds were randomly taken from each sealed plastic bags. Half of the seeds (30) were surface sterilized with 1% sodium hypochlorite (NaOCl) and the remaining (30 seeds) unsterilized. The surface treated and untreated seeds were plated on Potato Dextrose Agar (PDA), 10 seeds per Petri-plate and the petri-plates were incubated at 24 °C in alternating cycle of 12 hours light and 12 hours darkness for 7 days. The samples were incubated for additional 5 to 7 days to detect late appearing and slow sporulating fungi. Mounts from each fungal colony were prepared on a microscopic slide and then it was examined under high power magnification of a compound microscope. Fungi growing on seeds were identified up to species level by following standard manuals [3, 25 – 27]. The appearance of mycoflora and percentage occurrence of seed borne fungi detected from a total of 96 maize samples collected from four districts. The percentage of occurrence of fungi was calculated using the formula [22].

$$\text{occurrence of fungi (\%)} = \frac{\text{Number of seeds in which a particular fungus occurred}}{\text{total seed incubated}} \times 100$$

Method of data analysis

Statistical analysis was performed on response variables collected over the storage periods using ANOVA (Analysis of Variance) of SPSS (Statistical Package for the Social Sciences) Version 16.0. Means were compared for the significant factors by LSD (Least Significant Difference) test, and significance was accepted at 5%. Descriptive statistics was also used for organizing and presenting the data of the identified fungi.

RESULTS

Physical Quality Characteristics

Moisture content

At the beginning of the storage in the intermediate agro-ecology the average moisture content of the grain was 13.23% and 13.50% for the cobs in *Gombisa* and shelled maize grains in Sacks, respectively. The study, however, showed that, in the lowland agro-ecology, the average moisture content of the grain with the cob (*Gombisa*) and shelled grain (Sacks) was 14.58% and 13.58%, respectively (Figure 1). The average moisture content of maize just after harvest before storage was 13.37 ± 0.41 in intermediate and 14.08 ± 0.33 in lowland agro-ecologies. Grains stored in both storage containers lost moisture as storage time increased to 60 days reaching values of 10.07% and 10.57 % in *Gombisa* and Sack, respectively under intermediate agro-ecology. In the following storage days, grain with the cob stored in *Gombisa* continued to lose moisture attaining the lowest value of about 9.17% after 180 days. The situation with shelled grains stored in the Sacks was different in that the moisture

content exhibited increment to 11.08% at 120 days and 11.70% at 180 days of storage time. The grain moisture content recorded (Figure 1) both in *Gombisa* (14.58%) and sack (13.58%) under lowland agro-ecology during the initial loading day was found to be above the maximum safe moisture content (13.5%) for storing maize grain. This moisture content of maize grain is favorable for growth/ survival and multiplication of the fungal species identified in the study during 180 days of storage time. However, grain moisture content recorded in *Gombisa* (13.23%) and sack (13.50%) under intermediate agro-ecology is within the safe storage moisture content limit.

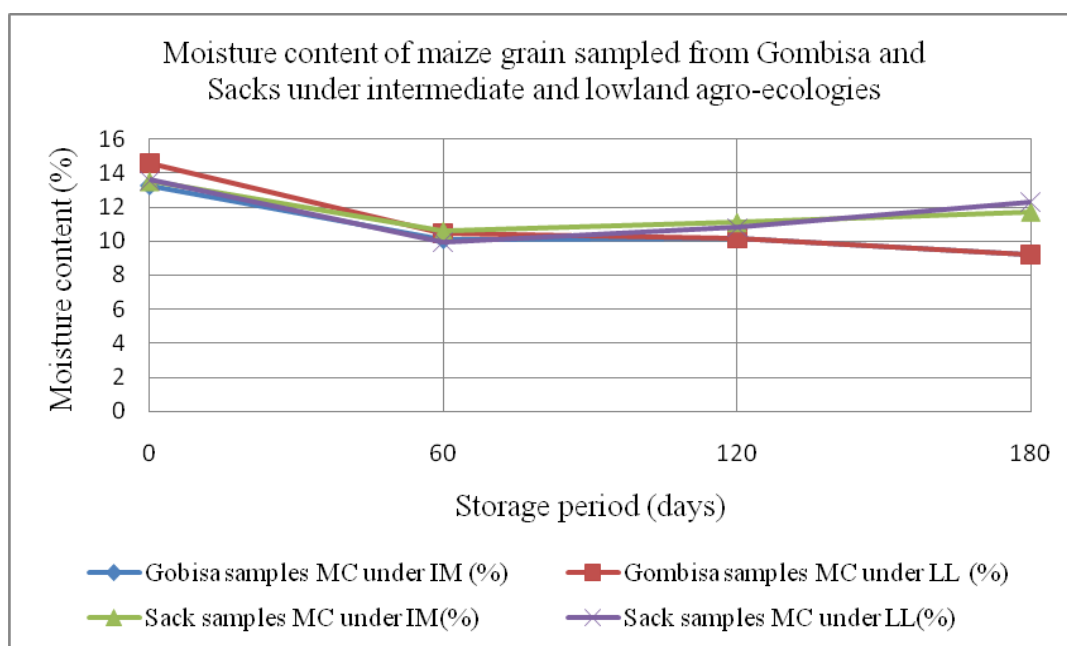


Figure 1: Moisture content (MC) of maize grain sampled from *Gombisa* and Sacks under intermediate (IM) and lowland (LL) agro-ecologies over storage periods of 180 days

Storage temperature and relative humidity

It is noted from Figures 2 and 3 that the average temperature had ranged from 18.50 to 30.00°C and 15.00 to 28.82 °C for *Gombisa* and Sack respectively under intermediate agro-ecology, while a temperature range of 21.30 to 35.00°C for *Gombisa* and 16.55 to 28.95°C for Sack were recorded with reference to lowland agro-ecologies. It is interesting to note that the average relative humidity under intermediate agro-ecology had ranged from 30.83 to 54.67% and 30.00 to 65.17% for *Gombisa* and Sack, respectively (Figure 2). Whereas the relative humidity ranges were 39.17 to 51.00% for *Gombisa* and 32.00 to 62.33% for Sack under lowland agro-ecology (Figure 3). These temperature and relative humidity records were optimal for the identified fungi species to flourish.

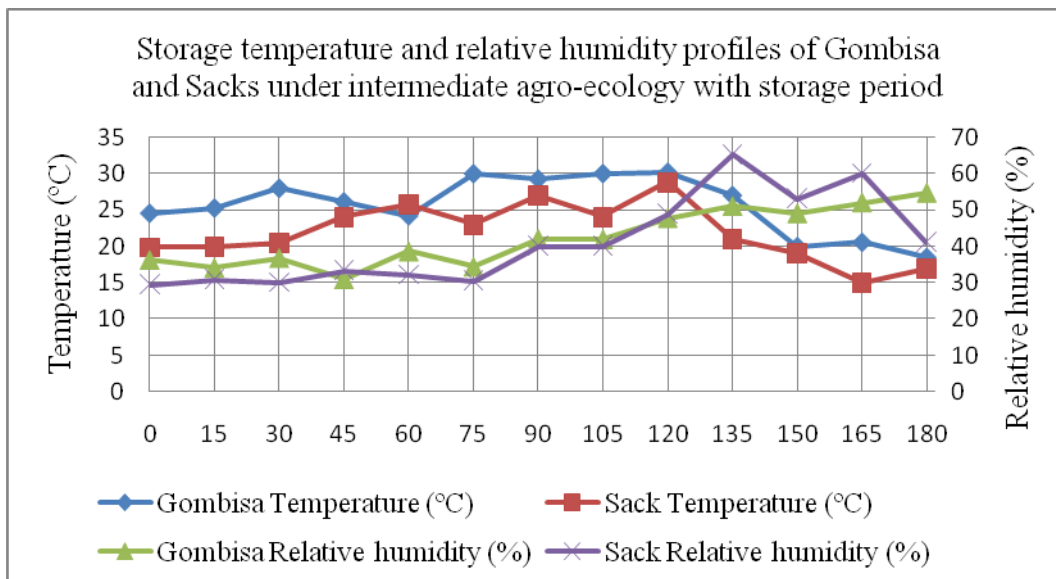


Figure 2: Storage temperature (°C) and relative humidity (%) profiles of Gombisa and Sacks under intermediate agro-ecology over storage periods of 180 days

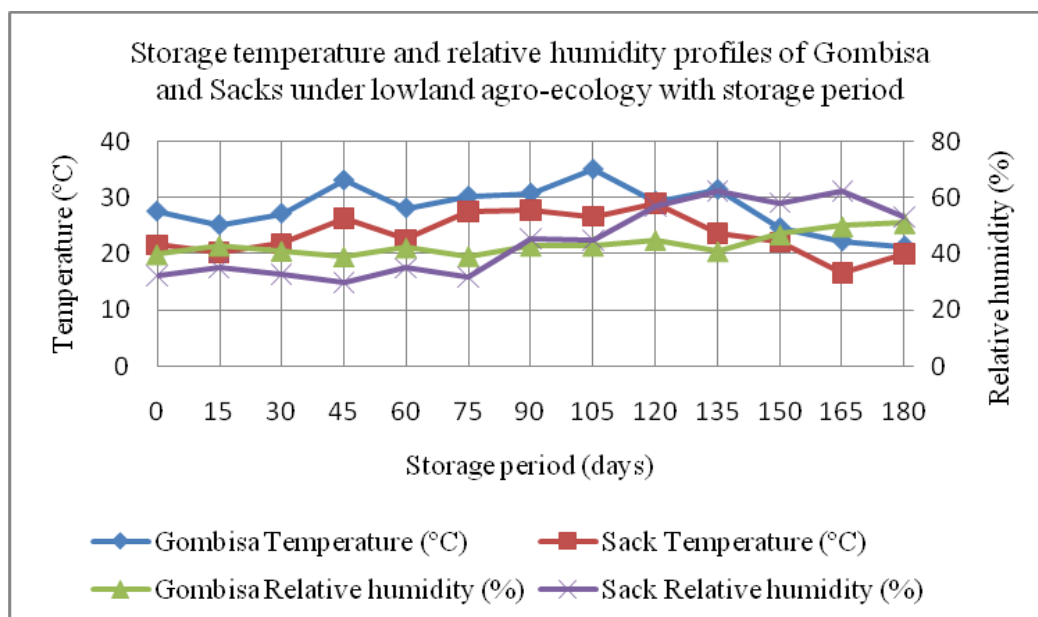


Figure 3: Storage temperature (°C) and relative humidity (%) profiles of Gombisa and Sacks under lowland agro-ecology with storage period

Effects of storage periods on viability of maize grain under intermediate and lowland agro-ecologies

The initial test for germination percentage under intermediate agro-ecology indicated 98% and 97.50 % for *Gombisa* and Sacks, respectively (Table 1). However, the values exhibited statistically significant ($P<0.05$) reduction as the storage period increased. After two months, it dropped to 87.5% for kernels collected from *Gombisa* while it was 92.0% in Sacks. There was a significant ($P<0.05$) reduction in germination percentage reaching 80.17% for grains collected from Sacks 120 days after storage. The germination capacity of kernels obtained from *Gombisa* further plunged to 68.50% in six months, while that of the kernels stored in the Sack dropped up to 120days.

Under lowland agro-ecologies 97.50% and 96.50 percentage of germination for *Gombisa* and Sacks respectively, was recorded during initial loading day. It was also noted that the values exhibited statistically significant ($P<0.05$) reduction as storage period increased to two months at which it dropped to 79.50% for kernels collected from *Gombisa* and 86.00% for kernels collected from Sacks. The results cited in Table 1 revealed that the values remain unchanged for the subsequent four and six months in both storage containers with final values of 70.17% and 83.83% in *Gombisa* and Sacks, respectively.

Identification of fungi

The observations in relation to fungal identification in the seeds of maize are given in Tables 2 and 3. All together eight species of fungi and one sterile white mycelium were isolated from maize seeds. These include *Aspergillus flavus* Link ex Grey, *Aspergillus fumigatus* Fres., *Aspergillus niger* Van Tieghem., *Aspergillus terreus*, *Cladosporium cladosporioides* (Fres) de Vries, *Drechslera halodes* Subram & Jain., *Fusarium oxysporum* Schlecht., Emend. Synd & Hans. and *Penicillium chrysogenum* Thom and (Wehmer) Westling.

The most common fungi isolated from grains stored in both storage containers (*Gombisa* and Sack) were *A. flavus*, *A. niger*, *D. halodes* and *F. oxysporum* in Kersa and Omo Nada of intermediate agro-ecology and in Sekoru and Tiro Afeta of lowland agro-ecology.

The results indicated that in almost all of the four districts these fungi were consistently recorded throughout 180 days of the storage period (Tables 2 and 3). In all of the studied districts, the presence of *A. fumigatus*, *C. cladosporioides*, *A. terreus* and *Penicillium spp.* were inconsistent with the increase in the storage days. During the course of the present study, the species *P. chrysogenum* and *A. terreus* were detected in districts found under intermediate agro-ecology and they were detected only from maize samples stored in *Gombisa* from Kersa and Omo Nada, respectively (Table 2).

It is also evident from the data in Table 4, that *A. flavus*, *A. niger*, *D. halodes*, and *F. oxysporum* were the most frequent species of fungi recorded over the storage periods

from 90%, 51%, 72% and 44% of the seed samples respectively, whereas *A. fumigatus* and *C. cladosporioides* were detected from 3.6 and 15% samples, respectively. Both *A. terreus* and *Penicillium* species were the most dominant species of fungi being detected only from 0.5% of the samples and sterile white mycelium was detected from 10% of seed samples.

DISCUSSION

It is clear from the results (Figure 1) that the differences in the initial grain moisture content of grains could be due to initial exposure of sampled grains to different ambient temperature and relative humidity. The loss of moisture after 180 days is probably due to the poor level of natural aeration in the grain stored in Sacks as compared to that in *Gombisa* along with the cumulative effect of respiration from the grain itself, insects and the mold population, which existed [28]. Further, polypropylene sack is moderately permeable to moisture, gases and odors [29]. The decrease in the moisture content from grains both in *Gombisa* and Sacks could be due to release of moisture in to the surrounding dry air. Interestingly, the decrease in the moisture content in *Gombisa* was higher than that in Sacks. This might be due to more ventilating capacity of the perforated wall of *Gombisa* than that of the Sack. In terms of moisture content the same trend was observed in both storage structures under lowland agro-ecology.

It was observed from the results (Figures 2 & 3) that the storage temperature and relative humidity recorded were optimal for the identified fungi species to flourish and cause maximum deterioration to maize grains.

It is evident from Table 1, that during the storage period, a gradual reduction in germination percentage of maize grain occurred for *Gombisa* under intermediate agro-ecologies, while the percentage was, relatively less, in case of Sacks. Germination loss in *Gombisa* and Sacks as the period of storage increased might be due to fungal invasion [30].

From Tables 2 and 3, it is concluded that, the most common fungi such as *A. flavus*, *A. niger*, *D. halodes* and *F. oxysporum* recorded in almost all the four districts throughout 6 months of storage period. While *Cladosporium cladosporioides* found upto 2 months for *Gombisa* in Kersa of intermediate agro-ecology and until 120 days in Sekoru of low land agro-ecology. *Penicillium chrysogenum* recorded for *Gombisa* after 4 months in OmoNada of Intermediate agro-ecology.

Presence of field fungal species like *C. cladosporioides* and *A. flavus* on maize stored in *Gombisa* and Sacks could be due to infestation of standing grain in the field, or from repeated use of the same drying floor year after year. The optimum temperature for growth of stored grain mold is about 25-30°C, but some species like *Aspergillus* spp. grow well at 35-37°C or above. Grains can be contaminated by spores of storage fungi during harvest, transport and handling operations [31]. A number of fungi isolated in the present study are known to produce mycotoxins which are harmful for

human health [11, 13]. Mycotoxins can cause severe damage to liver, kidney and nervous systems of human being even in low dosages [3]. *Fusarium* and *Aspergillus* species are common fungal contaminants of maize and also produce mycotoxins [4, 32]. *A. flavus* produces aflatoxins B₁, B₂, G₁ and G₂ which are carcinogenic and produce liver cancer [33-34]. Aflatoxins have powerful tetratogenic, mutagenic and hepato-carcinogenic effects [35]. *A. terreus* attacks human skin and nail and is parasitic on human ear [36]. *F. oxysporum* produce Zeralenone α and Zeralenone β that cause haemorrhage and necrosis in bone marrow. Other mycotoxins include trichothecenes and zearalenone, compounds known to injure the intestines, bone marrow, lymph nodes, spleen, and thymus. They are produced by species of *Fusarium* that grow on grain [33]. Besides, species of *Penicillium* cause kidney disease in human and pigs due to production of the mycotoxin called Citrinin [37].

CONCLUSION

The current study revealed that fungi such as *A. flavus*, *A. niger*, *D. halodes* and *F. oxysporum* are the major pathogens of maize stored in traditional storage structures (*Gombisa* and Sacks) under intermediate and low land agro-ecologies of Jimma zone, Ethiopia. As the storage period increased, the presence of some of the most common fungi pest were consistent, the damage they inflict to stored maize increases in both storage structures and ecologies affecting viability (germination capacity) of the seeds for the next planting season and the usability of the seeds as food and feed because of the potential toxins they may produce. This calls for future research in the area of investigating alternative eco-friendly fungi management methods such as the use of botanicals with fungicidal effects, physical and mechanical methods that modify the suitability of the physical environment for the multiplication of these fungi, identifying resistant and or tolerant maize varieties, identification/screening of effective bio-agents, screening of effective synthetic fungicides with novel chemistry. Furthermore, there is a strong need to train maize producers, traders and consumers who are involved in production and marketing chains of maize crop in the area of storage pests and their effective management. Improved storage structures need to be investigated for maize grain storage in the study area too.

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Table 1: Germination percentage of maize grain sampled from *Gombisa* and Sacks under intermediate and lowland agro-ecologies

Storage period (days)	Germination (%)*			
	Intermediate agro-ecology		Lowland agro-ecologies	
	<i>Gombisa</i>	Sacks	<i>Gombisa</i>	Sacks
Initial loading day	98.00 ± 0.52 ^a	97.50 ± 0.62 ^a	97.50 ± 0.76 ^a	96.50 ± 0.76 ^a
60	87.50 ± 1.23 ^b	92.00 ± 1.77 ^a	79.50 ± 7.29 ^b	86.00 ± 5.82 ^b
120	81.33 ± 4.73 ^b	80.17 ± 2.75 ^b	79.17 ± 4.24 ^b	83.67 ± 1.54 ^b
180	68.50 ± 2.90 ^c	80.50 ± 2.01 ^b	70.17 ± 4.29 ^b	83.83 ± 1.85 ^b
LSD (0.05)	10.50	11.50	18.00	10.50

*Means of twelve observations

Means ± standard error with different letters in a column are significantly different ($p < 0.05$) according to least significance difference test

Table 2: Appearance of mycoflora in Kersa and Omo Nada storage containers of intermediate agro-ecology

Fungi identified	Kersa								Omo Nada							
	Gombisa				Sack				Gombisa				Sack			
	ID	60	120	180	ID	60	120	180	ID	60	120	180	ID	60	120	180
<i>Aspergillus flavus</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
<i>A. fumigatus</i>	-	-	-	-	-	-	-	-	+	+	-	-	-	-	-	-
<i>A. niger</i>	+	+	+	+	+	+	+	+	+	+	+	-	+	+	+	+
<i>A. terreus</i>	-	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>Cladosporium</i>	+	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>cladosporioides</i>																
<i>Drechslera halodes</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
<i>Fusarium oxysporum</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
<i>Penicillium chrysogenum</i>	-	-	-	-	-	-	-	-	-	-	+	-	-	-	-	-
<i>Sterile mycelium (white)</i>	+	+	+	+	-	-	-	-	+	+	-	-	-	+	+	-

+ is the presence and - is the absence of fungi; ID, 60, 120 and 180 are storage periods in days; ID = Initial loading day

Table 3: Appearance of mycoflora in Sekoru and Tiro Afeta storage containers of lowland agro-ecology

Fungi identified	Sekoru								Tiro Afeta							
	Gombisa				Sack				Gombisa				Sack			
	ID	60	120	180	ID	60	120	180	ID	60	120	180	ID	60	120	180
<i>Aspergillus flavus</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
<i>A.fumigatus</i>	+	-	-	-	-	+	-	-	+	-	-	-	-	-	-	-
<i>A.niger</i>	+	+	+	-	+	+	+	+	+	+	+	+	+	+	+	+
<i>Cladosporium</i>	+	+	+	-	-	-	-	-	-	+	-	+	-	-	-	-
<i>cladosporioides</i>																
<i>Drechslera halodes</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+
<i>Fusarium oxysporum</i>	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+

+ is the presence and – is the absence of fungi; ID, 60, 120 and 180 are storage periods in days; ID = Initial loading day

Table 4: Percentage occurrence of fungi isolated from stored maize in selected districts of Jimma zone, Ethiopia

Mycoflora	Storage periods				%
	ID	60	120	180	
<i>Aspergillus flavus</i>	41	46	46	39	90
<i>A.fumigatus</i>	5	2	0	0	3.6
<i>A.niger</i>	25	29	25	18	51
<i>A.terreus</i>	0	1	0	0	0.5
<i>Cladosporium</i>					
<i>cladosporioides</i>	2	3	2	7	15
<i>Drechslera halodes</i>	33	41	37	27	72
<i>Fusarium oxysporum</i>	16	20	24	24	44
<i>Penicillium chrysogenum</i>	0	0	1	0	0.5
<i>Sterile mycelium (white)</i>	3	11	3	3	10

'0' indicates the absence of fungi; ID (Initial loading day), 60, 120 and 180 represent storage periods in days

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