

Bioconversion of Straw into Improved Fodder: Fungal Flora Decomposing Rice Straw

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The fungal flora decomposing rice straw were investigated all over the soil of Sharkia Province, east of Nile Delta, Egypt, using the nylon net bag technique. Sixty-four straw-decomposing species belonging to 30 genera were isolated by the dilution plate method in ground rice straw-Czapek's agar medium at pH 6. The plates were incubated separately at 5°C, 25°C and 45°C, respectively. Twenty nine species belonging to 14 genera were isolated at 5°C. The most frequent genus was *Penicillium* (seven species), and the next frequent genera were *Acremonium* (three species), *Fusarium* (three species), *Alternaria*, *Chaetomium*, *Cladosporium*, *Mucor*, *Stachybotrys* (two species) and *Rhizopus stolonifer*. At 25°C, 47 species belonging to 24 genera were isolated. The most frequent genus was *Aspergillus* (nine species), and the next frequent genera were ranked by *Penicillium* (five species), *Chaetomium* (three species), *Fusarium* (three species). Each of *Alternaria*, *Cladosporium*, *Mucor*, *Myrothecium* and *Trichoderma* was represented by two species. At 45°C, 15 species belonging to seven genera were isolated. These were seven species of *Aspergillus*, two species of *Chaetomium* and two species of *Emericella*, while *Hemicella*, *Malbranchea*, *Rhizomucor* and *Talaromyces* were represented by one species respectively. The total counts of fungi the genera, and species per gram of dry straw were significantly affected by incubation temperature and soil analysis ($P < 0.05$).

KEYWORDS: Bioconversion, Fodder, Fungal decomposition, Rice straw

Rice (*Oryza sativa* L), one of the world's leading crops, is cultivated in about 2×10^6 feddans in Egypt and the production of rice straw reaches about 8×10^6 tons per year. Straw is usually either burnt in the field causing environmental hazards such as respiratory diseases or disposed in a way that does not benefit the farmers to its maximum extent (Arai *et al.*, 1998; Samar *et al.*, 1999; Torigoe *et al.*, 2000). Fungi are the most important group among microbial agents for straw decomposition (Hudson, 1972; Srinivason, 1979; Harper and Lynch, 1982a; Yananobe *et al.*, 1994; Morais *et al.*, 1999; Tengerdy and Szakacs, 2003). There have been many surveys of cellulose-decomposing fungi but most fungi were isolated directly from soil or other sources on pure cellulose (Abdel-Hafez *et al.*, 1978; Abdel-Hafez and Abdel-Kader, 1980; Mazen *et al.*, 1980; Abdel-Hafez, 1982; Abdel-Kader *et al.*, 1983; Moubasher *et al.*, 1985). Surveys on wheat straw and other cultural wastes were conducted by other workers (El-Nawawy, 1972; El-Kady *et al.*, 1981; Abdel-Hafez *et al.*, 1990). Moubasher *et al.* (1985) isolated 19 species (one variety) belonging to 13 genera from wheat and broad bean straw compost at 45°C. However, few studies were focused on rice straw. Coronel *et al.* (1991) found that *Aspergillus fumigatus* was the most active cellulase producer on rice straw substrate among 144 tested strains of thermophilic lignocellulose-degrading fungi. Rai *et al.* (2001) isolated 42 mesophilic species from litter of rice

straw.

Research on the bioconversion of these agricultural residues into a microbial biomass as an improved feed supplement has been conducted in this study. The present investigation aimed the mycoflora inhabiting rice straw at 5°, 25° and 45°C in different localities.

Materials and Methods

Samples and soil analysis. Nylon net bag technique (House and Stinner, 1987; Wise and Schaefer, 1994) was used in this study. Thirty nylon bags each containing 10g of rice straw chopped to lengths of 5–6 cm, were prepared. These bags were distributed in different places throughout 10 areas (three bags in each area) of Sharkia Province of Egypt, in which large quantities of rice straw are produced. These sampling areas were; Abo-Kabir, Diarb-Negm, Hehia, Mashtool, Menia El-Kamh and Zagazig soils where were cultivated with *Triticum vulgare* and *Trifolium alexandrinum* after harvesting of paddy rice, *Oryza sativa*. Abo-Hamad and Fakoos areas were *Mangifera indica* orchards while Belbeis and Kafr-Sakr were orchards of *Citrus sinensis*. The bags were buried 5–10 cm from the soil surface for 10 days, then collected again for investigation.

Soil samples collected from the same localities at which nylon net bags were buried, were analyzed chemically for total soluble salts and organic matter content (Jackson, 1962). The soil texture was determined by the

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sieve method (Piper, 1947) using a standard, Rot-Top electric sieve shaker, VEB Metall Weberei Neustadt, Orla D.D.R. The reaction of the soil was measured using the glass electrode pH-meter (Richards, 1954).

Estimation of cellulose decomposing fungi. Fungi inhabiting rice straw were estimated using dilution plate method as described by Johnson *et al.* (1959). One ml of the straw suspension was introduced to each agar plate using Menzies dipper (1957) as recommended by Watson (1960) and Moubasher (1963). Modified Czapek's agar medium was used in which ground rice straw replaced sucrose (20 g/l) and to which rose Bengal (1/15000) was added as a bacteriostatic agent (Smith and Dawson, 1944), and the pH of the medium was adjusted to 6.0. Fifteen plates for each bag were used and incubated at 5°, 25° and 45°C (five plates at each temperature degree) for 15 days. The plates were examined daily. The average number of colonies per dish was multiplied by the dilution factor to obtain the number of colonies/g dry straw in the original rice straw sample.

Fungal identification. The fungal colonies appeared were picked up on Czapek's 0.5% yeast extract, malt extract and/or potato dextrose agar media, purified using spore technique and identified in this laboratory by consulting the following works: Gams *et al.*, 1998; Kubicek and Harman, 1998; Moubasher, 1993; Kitch and Pitt, 1992; Pitt, 1986; Domsch *et al.*, 1980; Pitt, 1979; Booth, 1977; Raper and Fennell, 1977; Ellis, 1971; Rifai, 1969; Raper and Thom, 1968 ; Cooney and Emerson, 1964.

Statistical analysis. The obtained data were conducted to one-way ANOVA and multiple-way ANOVA by Snedecor and Cochran (1982) and differences between means were done at the 5% probability level using Duncan's new multiple range tests (Duncan, 1955). Bivariate corre-

lation matrix of the obtained data was done using by SPSS software program (ver. 8) as described by Dytham (1999).

Results and Discussion

The straw dilutions (of each nylon net bag) used for estimation of rice straw-degrading fungi were 10^{-4} in the case of incubating the plates at 5° and 45°C, and 10^{-5} for 25°C. Rai *et al.* (2001) used dilution 10^{-4} for isolation of litter decomposing mycoflora of rice straw using nylon net bag technique. They also found that dilution plate method yielded maximum number of fungi in comparison with direct observation and damp chamber incubation methods.

Results in Tables 1 and 2 show that the total count of fungi per gram of dry straw was significantly affected ($P < 0.05$) by incubation temperature, pH value (–ve r), organic matter (+ve r) and total soluble salts (–ve r) of the soil where the nylon net bag was buried. The fungal-rich soils were sample no. 10, 9, 8, 4, 6 and 2, which are characterized by clay type, to be slightly acidic to neutral (6.9–7.2 pH-value) with high organic matter content (1.87–1.53%), and low total soluble salts (0.46–0.63%), and cultivated with *Triticum* and *Trifolium*. However the fungal-poor samples were soil sample no. 3 and 5 which are sandy, alkaline (7.8 and 7.7 pH), with low organic matter content (0.25 and 0.41%), high in total soluble salts (2.20 and 1.67%) and cultivated with *Citrus* and *Mangifera*. Abdel-Hafez (1978) and Helal (1993) found that the total population of fungi was influenced by the content of the total soluble salts of the soil. The highest fungal population was recorded from sample no. 10 (Zagazig area) which showed, 7.2×10^4 , 7.1×10^5 and 5.8×10^4 colony/g dry straw, while the lowest fungal population with 1.4×10^4 , 7.8×10^4 and 1.7×10^4 colony/g dry straw from sample number 3 (Belbeis area) at incubation temperature 5°, 25° and 45°C, respectively. Abdel-Sater and El-Said (2001)

Table 1. Characterization of soil samples tested and rice straw-decomposing fungi

Sample no.	Locality	Soil analysis				Total count/g dry rice straw			No. of genera			No. of species		
		Soil type	pH-value	OM%	TSS%	5°C	25°C	45°C	5°C	25°C	45°C	5°C	25°C	45°C
1	Abo-Hamad	Clay-sandy	7.5 b	0.85 c	0.91d	31297 ^x _d	236184 ^y _f	36184 ^x _{bc}	10 ^x _{bc}	15 ^y _c	4 ^x _b	21 ^x _{dc}	30 ^y _c	10 ^x _b
2	Abo-Kabir	Clay	7.2 c	1.53 b	0.63 e	52472 ^x _{bc}	453752 ^y _c	44793 ^x _{ab}	9 ^x _{cd}	17 ^y _d	4 ^x _b	20 ^x _e	33 ^y _{cd}	10 ^x _b
3	Belbeis	Sandy	7.8 a	0.25 e	2.20 a	13613 ^x _e	77952 ^y _i	16987 ^x _d	7 ^x _c	8 ^y _b	4 ^x _b	12 ^x _g	12 ^y _h	7 ^x _c
4	Diarb-Negm	Clay	6.9 cd	1.77 ab	0.53 ef	53520 ^x _{bc}	603839 ^y _c	54450 ^x _a	11 ^x _{ab}	21 ^y _b	6 ^x _a	24 ^x _{ab}	39 ^y _b	12 ^x _a
5	Fakoos	Sandy	7.7 ab	0.41 de	1.76 c	25713 ^x _{de}	221059 ^y _g	29668 ^x _{cd}	9 ^x _{cd}	12 ^y _g	4 ^x _b	15 ^x _f	22 ^y _g	8 ^x _c
6	Hehia	Clay	7.2 c	1.77 ab	0.60 ef	46422 ^x _c	524724 ^y _d	43979 ^x _{ab}	10 ^x _{bc}	18 ^y _c	5 ^x _{ab}	20 ^x _c	32 ^y _d	8 ^x _c
7	Kafir-Sakr	Clay-sandy	7.6 ab	0.62 cd	2.06 b	16754 ^x _{de}	186155 ^y _h	15241 ^x _d	8 ^x _d	13 ^y _f	4 ^x _b	14 ^x _f	25 ^y _f	7 ^x _c
8	Mashtool	Clay	7.1 c	1.77 ab	0.47 f	59104 ^x _{abc}	596859 ^y _c	54101 ^x _a	11 ^x _{ab}	18 ^y _c	5 ^x _{ab}	23 ^x _{bc}	34 ^y _c	12 ^x _a
9	Menia El-Kamh	Clay	7.0 cd	1.80 a	0.46 f	66434 ^x _{ab}	648051 ^y _b	54567 ^x _a	12 ^x _a	21 ^y _b	6 ^x _a	22 ^x _{cd}	41 ^y _a	12 ^x _a
10	Zagazig	Clay	6.8 d	1.87 a	0.46 f	72019 ^x _a	705051 ^y _a	58290 ^x _a	11 ^x _{ab}	23 ^y _a	6 ^x _a	25 ^x _a	42 ^y _a	12 ^x _a
	Mean					43677	425363	40838	10	17	5	20	31	10

1-The same letter in the same column/ row means not significant at $p < 0.05$.

2-a, b, c, ... for column comparison and x, y, z for row comparison.

Table 2. Pearson correlation matrix (r) and probability showing the relationship between soil analyses (pH, OM and TSS) and no. of genera, no. of species and total count at different incubation temperatures

Soil analysis	Total count			No. of genera			No. of species		
	Incubation temperature °C			Incubation temperature °C			Incubation temperature °C		
	5	25	45	5	25	45	5	25	45
OM	0.926**	0.956**	0.914**	0.783**	0.918**	0.607**	0.872**	0.898**	0.757**
PH	0.882**	0.903**	0.864**	0.724**	0.903**	0.596*	0.885**	0.885*	0.746**
TSS	0.842**	0.842**	0.842**	0.829**	0.893**	0.560*	0.899**	0.899**	0.812**

*Correlation is significant at the 0.01 level.

**Correlation is significant at the 0.001 level.

identified twenty three species belonging to 11 genera at 28°C from 30 samples of rice straw from Qena Governorate (South Valley), Egypt with total count 7.3×10^3 colonies/mg.

Sixty-four species belonging to thirteen genera were

collected in the present investigation (Table 3). The most diverse fungal flora was obtained at 25°C (24 genera and 47 species) with total fungal count 4.3×10^5 colony/g dry straw, followed by 14 genera and 29 species at 5°C with 3.6×10^4 colony/g dry straw, while only 7 genera and 15

Table 3. Characterization of rice straw-decomposing fungi isolated at 5°, 25° and 45°C in rice straw- Czapek's agar medium

Genus	Incubation temperature °C												TNS
	5				25				45				
	NS	NCI	OR	TC	NS	NCI	OR	TC	NS	NCI	OR	TC	
<i>Acremonium</i>	3	23	H	2815	1	3	R	1280	—	—	—	—	3
<i>Alternaria</i>	2	30	H	5433	2	30	H	20710	—	—	—	—	2
<i>Aspergillus</i>	—	—	—	—	9	30	H	141827	7	30	H	24584	10
<i>Botrytis</i>	1	1	R	23	—	—	—	—	—	—	—	—	1
<i>Botrytrichum</i>	—	—	—	—	1	15	M	11984	—	—	—	—	1
<i>Chaetomium</i>	2	27	H	3048	3	21	H	16638	2	21	H	1967	4
<i>Cladosporium</i>	2	30	H	6341	2	25	H	30134	—	—	—	—	2
<i>Cochliobolus</i>	—	—	—	—	2	11	M	6399	—	—	—	—	2
<i>Colletotrichum</i>	—	—	—	—	1	2	R	349	—	—	—	—	1
<i>Emericella</i>	—	—	—	—	2	13	M	6400	2	16	H	1257	2
<i>Epicoccum</i>	1	7	L	81	1	3	R	698	—	—	—	—	1
<i>Fusarium</i>	3	29	H	2827	3	26	H	33973	—	—	—	—	4
<i>Gliocladium</i>	—	—	—	—	1	12	M	2094	—	—	—	—	1
<i>Humicola</i>	—	—	—	—	1	9	M	3723	1	13	M	593	1
<i>Malbranchea</i>	—	—	—	—	—	—	—	—	1	3	R	70	1
<i>Mucor</i>	2	28	H	2641	2	26	H	28854	—	—	—	—	2
<i>Myrothecium</i>	—	—	—	—	2	22	H	24782	—	—	—	—	2
<i>Nigrospora</i>	1	5	L	81	1	3	R	698	—	—	—	—	1
<i>Oidiodendron</i>	1	1	R	12	—	—	—	—	—	—	—	—	1
<i>Penicillium</i>	7	30	H	5573	5	25	H	35369	—	—	—	—	9
<i>Phoma</i>	—	—	—	—	1	11	M	3839	—	—	—	—	1
<i>Pestalotia</i>	1	1	R	23	—	—	—	—	—	—	—	—	1
<i>Rhizomucor</i>	—	—	—	—	—	—	—	—	1	30	H	11914	1
<i>Rhizopus</i>	1	28	H	1827	1	11	M	7330	—	—	—	—	1
<i>Scopulariopsis</i>	—	—	—	—	1	3	R	582	—	—	—	—	1
<i>Stachybotrys</i>	2	30	H	5376	1	13	M	5468	—	—	—	—	2
<i>Talaromyces</i>	—	—	—	—	—	—	—	—	1	10	M	314	1
<i>Trichoderma</i>	—	—	—	—	2	26	H	36300	—	—	—	—	2
<i>Trichothecium</i>	—	—	—	—	1	9	M	5236	—	—	—	—	1
<i>Verticillium</i>	—	—	—	—	1	2	R	233	—	—	—	—	1
No. of genera	14				24				7				64
No. of species	29		36101		47		424900		15		40699		

NS : number of species, NCI : number of cases of isolation, OR : occurrence remarks, TC : total count, TNS : total number of species.

H=high occurrence isolated more than 15 cases (out of 30).

M= moderate occurrence from 8 to 15 cases.

L= low occurrence from 4 to 7 cases.

R= rare occurrence, less than 4 cases.

Aspergillus represented 100% of the samples at 25° and 45°C, constituting 33.4% and 60.4% of total fungi, respectively, but did not appear at 5°C. *A. awamori*, *A. flavus*, *A. fumigatus*, *A. niger* and *A. terreus* were of high occurrence at both 25°C and 45°C, while *A. flavus* var *columinaris* was of low occurrence at both temperatures. At

Chaetomium came third after *Aspergillus* and *Penicillium*. It was isolated at 5°, 25° and 45°C and represented 90, 73 and 73% of the samples constituting 8.4, 3.9 and 4.8% of total count of fungi, respectively. *C. cochliodes* was of moderate occurrence and *C. globosum* was of high occurrence at 5° and 25°C, *C. spirale* was of moderate occurrence at 25° and 45°, while *C. thermophilum*, a thermophilic fungus, occurred only at 45°C. Soyong (1991) found that some *Chaetomium* species isolated from the

[illegible]

Table 4. Continued

Species	Incubation temperature °C								
	5			25			45		
	TC	NCI	OR	TC	NCI	OR	TC	NCI	OR
<i>Botrytis cinerea</i> Pers.	23	1	R	–	–	–	–	–	–
<i>Botrytrichum piluliferum</i> Sacc. & Marchal	–	–	–	11984	15	M	–	–	–
<i>Chaetomium cochliodes</i> Pall.	768	15	M	1862	8	M	–	–	–
<i>C. globosum</i> Kunze ex Stend.	2280	27	H	8959	18	H	–	–	–
<i>C. spirale</i> Zopf	–	–	–	5817	14	M	1164	13	M
<i>C. thermophilum</i> La Touche	–	–	–	–	–	–	803	9	M
<i>Cladosporium cladosporioides</i> (Fresen.) de Vries	6341	30	H	10239	14	M	–	–	–
<i>C. herbarum</i> (Pers.) Link	7574	30	H	19895	23	H	–	–	–
<i>Cochliobolus lunatus</i> Nelson & Haasis	–	–	–	4421	10	M	–	–	–
<i>C. sativus</i> (Ito & Kurib.) Drechsler ex Dastur	–	–	–	1978	5	L	–	–	–
<i>Colletotrichum dematium</i> (Pers. ex Fr.) Grove	–	–	–	349	2	R	–	–	–
<i>Emericella nidulans</i> (Eidam) Vuillemin var. <i>Lata</i> Subramanian	–	–	–	5236	12	M	1001	14	M
<i>E. quadrilineata</i> (Thom & Raper) Benj.	–	–	–	1164	5	L	256	7	L
<i>Epicoccum nigrum</i> Link	81	7	L	698	3	R	–	–	–
<i>Fusarium moniliforme</i> Sheld.	256	12	M	2792	7	M	–	–	–
<i>F. oxysporum</i> Schlecht	–	–	–	19197	24	H	–	–	–
<i>F. pallidoroseum</i> (Cooke) Sacc.	442	17	H	–	–	–	–	–	–
<i>F. solani</i> (Mart.) Sacc.	2129	28	H	11984	17	H	–	–	–
<i>Gliocladium catenulatum</i> Gilm. & Abbott	–	–	–	2094	12	M	–	–	–
<i>Humicola grisea</i> Traaen var. <i>grisea</i>	–	–	–	3723	9	M	–	–	–
<i>H. grisea</i> var. <i>thermoidea</i> Cooney & Emers.	–	–	–	–	–	–	593	13	M
<i>Malbranchea sulfurea</i> (Miche) Sigler & Carmich.	–	–	–	–	–	–	70	3	R
<i>Mucor circinelloides</i> Van Tiegh.	419	13	M	3490	11	M	–	–	–
<i>M. racemosus</i> Fresen.	2222	28	H	25364	25	H	–	–	–
<i>Myrothecium roridum</i> Tode	–	–	–	22920	22	H	–	–	–
<i>M. verrucaria</i> (Alb. & Schwein.) Ditmar	–	–	–	1862	5	L	–	–	–
<i>Nigrospora sphaerica</i> (Sacc.) Mason	81	5	L	698	3	R	–	–	–
<i>Oidiodendron griseum</i> Robak	12	1	R	–	–	–	–	–	–
<i>Penicillium canescens</i> Sopp	128	7	L	–	–	–	–	–	–
<i>P. chrysogenum</i> Thom	1757	27	H	–	–	–	–	–	–
<i>P. citrinum</i> Thom	791	19	H	15241	21	H	–	–	–
<i>P. corylophilum</i> Dierckx	163	7	L	4072	9	M	–	–	–
<i>P. herquei</i> Bain. & Sart.	23	2	R	–	–	–	–	–	–
<i>P. janthinellum</i> Biourge	–	–	–	233	1	R	–	–	–
<i>P. oxalicum</i> Currie & Thom	2315	29	H	15125	20	H	–	–	–
<i>P. rubrum</i> Stoll	–	–	–	698	3	R	–	–	–
<i>P. verrucosum</i> Dierckx var. <i>verrucosum</i>	396	14	M	–	–	–	–	–	–
<i>Phoma herburum</i> Peyronel	–	–	–	3839	11	M	–	–	–
<i>Pestalotia</i> sp.	23	1	R	–	–	–	–	–	–
<i>Rhizomucor pusillus</i> (Lindt) Schipper	–	–	–	–	–	–	11914	30	H
<i>Rhizopus stolonifer</i> (Lindt) Schipper	1827	28	H	7330	11	M	–	–	–
<i>Scopulariopsis brevicaulis</i> (Sacc.) Bain.	–	–	–	582	3	R	–	–	–
<i>Stachybotrys chartarum</i> (Ehrenb. ex Link) Hugh.	3165	29	H	5468	13	M	–	–	–
<i>S. elegans</i> (Pidopl.) Gams.	2211	30	H	–	–	–	–	–	–
<i>Talaromyces thermophilus</i> Stolk	–	–	–	–	–	–	314	10	M
<i>Trichoderma harzianum</i> Rifai	–	–	–	13031	19	H	–	–	–
<i>T. koningii</i> Oudem.	–	–	–	23269	26	H	–	–	–
<i>Trichothecium roseum</i> (Pers.) Link ex Gray	–	–	–	5236	9	M	–	–	–
<i>Verticillium catenulatum</i> (Kamyschko ex Barron & Onions) Gams	–	–	–	233	2	R	–	–	–
Sterile mycelia	–	–	–	465	3	R	–	–	–

TC : total count, NCI : number of cases of isolation, OR : occurrence remark.

H = high occurrence isolated more than 15 cases (out of 30).

M = moderate occurrence from 8 to 15 cases.

L = low occurrence from 4 to 7 cases.

R = rare occurrence, less than 4 cases.

rhizosphere of some economical plants can degrade rice husks, rice straw and paper.

Fusarium occupied the fourth place was isolated at 5° and 25°C but did not appear at 45°C and represented 96.7 and 86.7% at both temperatures constituting 7.8 and 8.0% of the total count of fungi, respectively. *F. moniliforme* was of moderate occurrence and *F. solani* was of high occurrence at both 5° and 25°C. *F. oxysporum* was of high occurrence at 25°C only, while *F. pallidoroseum* (*F. semitectum*) was of high occurrence at 5°C. El-Kady *et al.*, (1981) isolated *F. oxysporum* and *F. moniliforme* in high occurrence on cellulose medium from wheat straw. Also, Rai *et al.* (2001) reported that *F. semitectum* was dominant on rice straw.

Acremonium consisted of three species. *A. fusidioides* was of moderate occurrence and *A. kiliense* was of high occurrence only at 5°C. *A. strictum* appeared at 5 and 25°C in low and rare occurrence, respectively. *Acremonium* represented 76.7 and 10% of the samples constituting 7.8 and 0.3 of the total count at 5° and 25°C.

Cladosporium was isolated only at 5° and 25°C, and represented 100% and 83.3% of the samples constituting 17.6 and 7.1 of the total count of fungi. *C. cladosporioides* was of high occurrence at 5°C and moderate occurrence at 25°C, while *C. herbarum* was of high occurrence at both temperatures.

Cochliobolus appeared only at 25°C, represented in 36.7% of the samples constituting 1.5% of the total count. *C. lunatus* was of moderate occurrence, while *C. sativus* was of low occurrence.

Humicola appeared at 25° and 45°C represented 30 and 43.3% of the samples, constituting 0.9 and 1.5% of the total count of fungi. *H. grisea* Traaen was of moderate occurrence at 25°C only, while *H. grisea* var *thermoidea* (*Scytalidium thermophilum*) was of moderate occurrence only at 45°C.

Emericella nidulans (*Aspergillus nidulans*) was isolated at 25° and 45°C and represented 43.3 and 53.3% of the samples constituting 1.5 and 3.1% of the total fungi. *E. nidulans* var. *Lata* was of moderate occurrence at 25 and 45°C and *E. quadrilineata* occurred low at both temperatures.

Mucor was isolated at 5° and 25°C, represented 93 and 86.7% of the samples constituting 7.3 and 6.8% of the total fungi. *M. circinelloides* was of moderate occurrence and *M. racemosus* was of high occurrence at both temperatures.

Myrothecium was isolated only at 25°C, represented 73.3% of the samples constituting 5.8 of the total fungi. *M. roridum* was of high occurrence, while *M. verrucaria* was of low occurrence.

Stachybotrys was isolated at 5° and 25°C, represented 100% and 43.3% of the samples constituting 14.9 and 1.3% of the total count of fungi. *S. chartarum* was of high

occurrence at 5°C and moderate occurrence at 25°C, while *S. elegans* was of high occurrence only at 5°C.

Trichoderma appeared only at 25°C and represented 86.7% of the samples constituting 8.5% of the total fungi. This genus consisted of two species *T. harzianum* and *T. koningii* which were of high occurrence. *Trichoderma* species were often cited as high cellulose-decomposers (Moharram *et al.*, 1995; Tengerdy and Szakacs, 2003; Zayed and Abdel-Motaal, 2005).

The other genera that appeared in this study represented one species. *Botrytis cinerea*, *Oidiodendron griseum* and *Pestalotia* sp. appeared only at 5°C. *Botrytrichum piluliferum*, *Colletotrichum dematium*, *Gliocladium catenulatum*, *Phoma herburum*, *Scopulariopsis brevicaulis*, *Trichothecium roseum* and *Verticillium catenulatum* appeared only at 25°C. *Malbranchea sulfurea*, *Rhizomucor pusillus* and *Talaromyces thermophilus* appeared only at 45°C. *Epicoccum nigrum*, *Nigrospora sphaerica* and *Rhizopus stolonifer* appeared at 5° and 25°C. No species appeared at the three incubation temperatures, 5°, 25° and 45°C. Most of these fungi isolated on rice straw during this study were isolated with different frequency of occurrence from various hemicellulosic and cellulosic materials in Egypt (Moubasher and Mazan, 1990; Moubasher, 1993; Abdel-Sater and El-Said, 2001) as well as other parts of the world (Caldwell, 1973, Bisaria and Ghose, 1981; Maheshwari *et al.*, 2000).

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