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# Seed Borne Mycoflora of Tribal farmers' Saved Hill Rice, *Oryza sativa* in Northeast of India

A. Ratankumar Singh<sup>1</sup> • T. Boopathi<sup>1</sup> • S. B. Singh<sup>1</sup> • S. K. Dutta<sup>1</sup> • L. S. Singh<sup>2</sup> • Lungmuana<sup>1</sup> • S. Saha<sup>1</sup> • N. H. Singh<sup>1</sup>

<sup>1</sup>ICAR Research Complex for North Eastern Hill Region, Mizoram Centre, Kolasib-796081, Mizoram

<sup>2</sup>Department of Plant Pathology, Bidhan Chandra Krishi Viswavidyalaya, Mohanpur, Nadia- 741252, West Bengal.

# ARTICLE INFO

#### ABSTRACT

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Detection of seed borne pathogens through seed health testing is a critical step in the management of diseases for healthy crop establishment. The tribal farmers of Northeastern India saved their own rice seed which remains poor quality and act as carrier of many mycoflora that hinder seed viability and crop stand. This study was carried out to assess seeds of tribal farmers' saved hill rice and improved varieties (twenty local landraces and four improved varieties) for their seed borne mycoflora collected from Mizoram, Northeastern India. The seeds were subjected to blotter and agar plate techniques to identify various seed borne mycoflora and their germination per cent and seed vigour index also calculated by paper towel method. A total of 21 fungi were recorded like Acremoniella sp. Alternaria tenuis, Curvularia lunata, C. oryzae, Dreschslera oryzae, Fusarium moniliforme, F. oxysporum, F. semitectum, Microdochium oryzae, Pyricularia grisea, Rhizoctonia sp., Saracladium oryzae, Tilletia sp. Trichothesiums sp., stilaginoidea virens, Aspergillus flavus, A. niger, Penicillium sp. and Rizopus stolonifer and among these pathogen, Fusarium moniliforme was pre dominant in all tested rice samples ranging from (15-35%) and (54-82.0%) of agar plate and blotter paper method, respectively. Maipum (1317.43) and Manipur Nem (1171.76) showed the better performance in terms of seed germination and seedling vigour index. Idaw, recorded the maximum number of pathogen genera (14), least germination (45.45%) and vigour index (37.42). Tribal famers' saved hill rice seeds are found to be more associated with seed borne mycoflora than the improved varieties.

# 1. Introduction

Rice (*Oryza sativa* L.) is the main staple food of the North Eastern Region (NER) of India and West Bengal, Uttar Pradesh, Madhya Pradesh, Punjab, Orissa and Bihar are the major rice producing states. The NER comprised of seven mountainous states (Manipur, Mizoram, Nagaland, Tripura, Meghalaya, Arunachal Pradesh and Sikkim) of Indian Territory forming 7.8% of the total land area and share about 4% of the total population of the country which dominated by tribes. In NE region, rice is cultivated at hilly agro-ecosystem, occupying 3.51 million hectares which accounts for more than 80% of the total cultivated area of the region and 7.8 per cent of the total rice area in India. The total rice production of NER is estimated to be around 5.50 million tones with average productivity of 1.57 t/ha, which is much below the national average of 2.08 t/ha. The reason for such low productivity are several constraints in the production of rice, of which seed borne diseases caused by bacteria, fungi, viruses and nematodes are responsible for major economic losses in north east India. Agrawal (1999) reported that more than 50 fungal pathogens found to be seed-borne and it's deteriorated both quantity and quality aspects of rice (Janardhana et al. 1998; Kavitha et al. 2005). Rice seeds are infected by large number of fungi and perpetuated from one

<sup>\*</sup>Corresponding author: ratanplantpatho@gmail.com

season to another through infected seeds (Zope and Thrimurty 2004). Moreover, in this region, high rainfall and humidity during Kharif season exposed paddy seeds to many fungal invasions (Islam and Borthakur 2012). In India, total seed requirement is met up 20 % by certified seed and remaining 80% from farmers saved seed (Raj et al. 2007; Atwal 2013). Despite of availability of certified seeds, traditionally, tribal farmer of this region continue to produce their own local rice seed and reuse it without knowing the health status of seeds. Saved rice seeds were stored in very unhygienic conditions; hence it highly prone to seed inhabiting mycoflora which are capable of deteriorating seed quality and poor stands. In changing climate, many minor seed borne pathogens like false smut, leaf scald may act major pathogens and impart threat to tribal hill resource poor farmers. Moreover, no research has

been done in Mizoram which sharing common agricultural practices with others NE states to evaluate health status of landraces rice seed from both authorized and unauthorized seed sectors. Since, every seed could play a vital role in the development of epidemics in fields, good quality and healthy seed of rice should be made available to farmers in order secure their production in increasing population. Several fungal pathogens have been isolated from rice grains and have been reported to be responsible for a number of diseases from the nursery to the field (Ibiam et al. 2006). Considering the above facts, the present study were undertaken to visualize different seed borne mycoflora and incidence of tribal farmers' saved hill rice seed collected from different regions of Mizoram, North eastern India. The result provides a database for further study to develop an effective management strategy of the pathogens.

**Table 1.** Details information of the tribal farmer saved hill rice seed samples collected from different hilly locations of Mizoram, North eastern region of India

S.	Name of landraces/	Habitat	Seed colour and	Sources of Collection						
No.	varieties	Туре	characteristics							
1	Idaw	Jhum	Light brown	Farmer-saved seed-Mizoram						
2	Akbuh	Jhum	Dark red with long awn	Farmer-saved seed-Mizoram						
3	Zakew	Jhum	Light red with long awn	Farmer-saved seed-Mizoram						
4	Saii Buh	Jhum	Light brown with long awn	Farmer-saved seed-Mizoram						
5	Zaitlai	Jhum	Light brown	Farmer-saved seed-Mizoram						
6	Vai buh	Jhum	Light brown	Farmer-saved seed-Mizoram						
7	Zaizpuii	Jhum	Light brown with long awn	Farmer-saved seed-Mizoram						
8	Fazai	Jhum	Light brown	Farmer-saved seed-Mizoram						
9	Manipur (Rum)	Lowland	Light brown	Farmer-saved seed-Mizoram						
10	Vuitawi	Lowland	Light brown	Farmer-saved seed-Mizoram						
11	Buh tawi sang	Lowland	Light brown	Farmer-saved seed-Mizoram						
12	Buh Mui	Lowland	Reddish	Farmer-saved seed-Mizoram						
13	Shan buh	Lowland	Light brown	Farmer-saved seed-Mizoram						
14	Sawkar Buh	Lowland	Light brown	Farmer-saved seed-Mizoram						
15	Thlarau Buh	Lowland	Light brown	Farmer-saved seed-Mizoram						
16	Zoro	Lowland	Light brown	Farmer-saved seed-Mizoram						
17	Thingtlang vai Buh	Lowland	Light brown	Farmer-saved seed-Mizoram						
18	Tauphai Buh	Lowland	Light brown	Farmer-saved seed-Mizoram						
19	Manipur (Nem)	Lowland	Light brown	Farmer-saved seed-Mizoram						
20	Maipuam	Lowland	Light brown	Farmer-saved seed-Mizoram						
21	RC Maniphou-9	Lowland	Light brown	ICAR-RC-NEH Region, Manipur Centre,						
				Imphal, Manipur, India						
22	RC Maniphou-10	Lowland	Light brown	ICAR-RC-NEH Region, Manipur Centre,						
				Imphal, Manipur, India						
23	RC Maniphou-11	Lowland	Light brown	ICAR-RC-NEH Region, Manipur Centre,						
				Imphal, Manipur, India						
24	CAU-R-1(Tampha	Lowland	Light Brown	Central Agricultural University, Imphal,						
	phou)			Manipur, India						

## 2. Materials and Methods

#### 2.1 Paddy seed sample collection

The study area is situated in Himalayan hill range, Mizoram, North-East India (23°28'40"N and 93°19' 44''E) with an average altitude of 1678 m (MSL). In this region, the traditional landraces of rice is widely cultivated due to its suite to food habit, high palatability (sticky) and adaptability in the prevailing climatic conditions. Topographically, maximum areas under rice are grown in Jhum (shifting cultivation) and valley (low land). Twenty landraces seed samples and four improved varieties (approximately 2 kg) (Table 1) were collected from different the hill/Jhum farmers of Mizoram during harvest season of 2012-2014. Seed samples were brought to the laboratory in sterile plastic bag and kept at 4°C until the diagnosis of pathogens. All seed samples were subjected to seed health testing using blotter technique, agar plate technique, (ISTA 1999) germination by paper towel method and vigour index was evaluated based on seedling length (ISTA 1999; Abdul Baki and Anderson (1972).

#### 2.2 Detection of seed mycoflora

Seed borne fungi were detected by using the blotter test method developed by (ISTA 1999; Mathur and Konssdal 2003). Four hundred seeds were randomly selected from each sample and placed on three layers of moisten sterilized blotter paper at the rate of 20 seeds/ Petri plate (90 mm dia.). The seed plated were incubated at 22±1°C in an incubator for 10 days maintaining 12 h alternate cycles of light and darkness. After incubation, fungi associated with seeds were isolated by pure culture method and examined under different magnification of compound and stereomicroscope for presence of mycoflora. Identification of isolated mycoflora was done based on their morphological characters and their microscopic examination of spores with help of available literatures (Barnett and Hunter 1972; Mathur and Kongsdal 2003; Mew and Gonzales 2003). The percent incidence of the seed mycoflora was recorded in each sample and the data were tabulated for statistical analysis.

# 2.3 Agar plate method

Another set of experiment was also carried out on agar plate technique; four hundred seeds were tested for each sample maintaining twenty seeds per plate with 20 replications. The plated seeds were incubated for 5 days  $22\pm1^{\circ}$ C under 12h altering cycles of light and darkness. At the end of the incubation period, fungi growing out from the seeds on the agar medium were kept under constant examination and identification was done as mentioned above.

## 2.4 Testing of seed germination and seedling vigour

The method developed by Warham (1990) was followed. Three replicates of 100 seeds each were incubated in wet blotter paper towels for a period of 15 days for germination test according to ISTA under standard conditions of light, temperature and humidity. The paper towels were rolled and the ends were closed by rubber band and covered by butter paper to prevent drying up. For determination of seedlings vigour, randomly ten seedlings were selected from each paper and their individual shoot and root length was measured. Shoot length (cm) was measured from the base of the stem up to the growing point of the youngest leaf. Similarly, root length (cm) was also measured from the shoot and root juncture point to the largest available lateral root apex. The vigour index of the seedlings was calculated using following formula developed by Abdul Baki and Anderson (1972). Germination (%) = (Number of seeds germinated/ Total number of seeds tested) × 100

Vigour index = (Mean of root length + Mean of shoot length) × Percentage of seed germination

The laboratory experiment was conducted following Completely Randomized Design (CRD) and recorded data on various parameters under the present study were statistically analyzed using SAS Software Version 9.3 (SAS Institute Inc. 2011).

#### 3. Results and Discussion

In blotter technique, a total of 19 seed borne mycoflora were recorded like, Acremoniella sp. Alternaria tenuis, Curvularia lunata, C. oryzae, Dreschslera oryzae, Fusarium moniliforme, F. oxysporum, F. semitectum, Microdochium oryzae, Pyricularia grisea, Rhizoctonia sp., Saracladium oryzae, Tilletia sp., Trichothesiums sp., Ustilaginoidea virens, Aspergillus flavus, A. niger, Penicillium sp. and Rizopus stolonifer (Table 2). Among the pathogens, Fusarium moniliforme infected the highest incidence (15-35%) on all rice seeds tested whereas, Saii Buh and Idaw, jhum rice recorded the maximum incidence of 35% and 30%, respectively. Improve rice varieties recorded less pathogen load as compared to jhum and lowland landrace rice seed. Alternaria tenuis, Pyricularia grisea, Rhizoctonia sp., Saradocladium oryzae, Tilletia sp., Ustilagoinedea virens and Fusarium semectum were not frequently occurred on all rice samples whereas Fusarium moniliforme, Ustilaginoidea virens, Aspergillus flavus, A. niger, Penicilium sp. and Rhizopus stolonifer were predominant seed mycoflora on seed tested. Among the different landraces rice, Maipum

Pathogens	% incidence seed borne mycoflora																							
															Improv	Improve cultivars								
	Idaw	Ak buh	Zakew	Saii Buh	Zaitlai	Vai buh	Zaipui	Fazai	M (Rum)	Vuitawi	Buh tawi sang	Buh Mui	Shan buh	Sawkar Buh	Thlarau Buh	Zoro	Thingtlang vai Buh	Tauphai Buh	M (Nem)	laipum	RCM- 9	RCM- 10	RCM- 11	CAU R-1
Acremoniella sp.	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	2.50	0.00	3.50	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	2.00	2.20	0.10	0.20
Alternaria tenuis	3.50	0.00	0.00	0.00	7.50	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.50	0.00	0.00	0.00	0.00	0.00	0.00
Curvularia lunata	10.0	12.0	9.00	15.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	10.0	19.0	10.0	10.0	0.00	7.00	0.00	0.00	3.20	2.50	0.30	2.30
Curvularia oryzae	0.50	0.10	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.00	3.00	3.50	3.50	0.00	1.00	0.00	0.00	0.00	0.00	0.00	0.00
Dreschslera oryzae	0.00	0.00	0.00	3.00	0.00	4.00	0.00	0.00	0.00	0.00	0.00	2.0	0.00	1.00	20.0	0.00	0.00	0.00	0.00	2.30	0.00	0.00	0.00	0.00
Fusarium moniforme	30.0	17.0	25.0	35.0	21.0	30.0	15.0	15.5	26.0	15.0	10.0	30.0	15.0	10.00	16.0	16.0	10.0	19.0	10.0	16.3	0.30	1.50	10.0	4.25
Fusarium oxysporum	9.00	10.0	3.00	0.50	4.00	6.00	7.50	0.00	10.0	3.00	0.00	10.3	11.0	3.00	0.00	0.00	1.50	7.00	3.50	0.00	1.75	0.75	0.50	5.00
Fusarium semitectum	0.9	0.00	0.00	0.00	0.00	0.00	0.00	1.50	0.10	0.00	0.00	0.00	0.00	0.00	3.50	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Microdochium oryzae	3.50	0.50	7.50	3.75	10.25	3.75	6.00	12.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Pyricularia grisea	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	2.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Rhizoctonia sp.	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.75	0.00	0.00	0.00	0.00	4.31	0.00	0.00	0.00	0.00
Saracladium oryzae	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.25	0.75	1.25	0.25	0.00	4.25	0.00	0.25	0.25	1.75	3.75	0.00	0.00	0.00	0.00	0.00
Tilletia sp.	0.00	0.00	0.00	0.00	0.00	0.00	3.00	5.50	0.00	0.00	0.16	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Trichothesiums sp.	0.00	0.00	0.00	0.00	0.00	0.00	2.00	0.00	0.00	0.00	0.10	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Ustilaginoidea virens	2.10	5.00	6.00	4.00	8.00	10.00	3.00	5.00	6.00	5.00	6.00	8.00	8.00	9.00	10.0	8.00	12.0	14.0	12.5	0.00	5.60	1.40	2.50	3.20
Aspergillus flavus	0.10	0.15	0.10	2.75	0.00	1.00	3.00	0.30	0.90	3.00	0.10	5.50	3.60	3.00	5.00	3.00	4.00	5.00	3.00	0.00	5.00	2.00	3.00	1.00
Aspergillus niger	3.50	7.00	7.00	3.00	9.00	8.00	7.00	6.00	1.00	0.00	10.0	10.0	6.50	10.30	5.00	2.50	3.50	3.25	0.75	0.56	3.15	1.75	1.00	7.25
Penicillium sp.	13.0	10.0	14.0	3.25	10.0	14.0	15.0	13.0	9.00	11.0	11.0	8.30	10.50	10.0	5.50	18.0	16.00	13.00	12.00	11.0	00.50	10.50	11.00	12.50
Rizopus stolonifer	5.00	3.00	0.00	3.00	0.00	5.00	3.00	0.00	5.00	3.00	0.00	1.00	0.00	1.00	2.00	1.00	0.50	0.10	0.00	0.00	3.00	1.00	3.00	3.00
Nos .of Genera/species	12	10	8	10	7	9	10	8	10	9	9	9	7	11	11	9	8	11	7	5	9	9	8	9

Table 2. Percent incidence of different seed borne mycoflora from farmer-saved rice seed of Mizoram, NE Region of India by blotter method

M=Manipur, RCM=Research Centre Maniphou, CAU-R=Central Agricultural University Rice

recorded the least number of pathogenic genera (5), followed by Zatlai (7), Buhmuii (7), Fazai (8) and Zakew (8). Authentic literature on seed borne mycoflora of tribal farmer saved hill rice seed is limited, but some researchers reported on improved rice cultivars. Sharma (1987) detected 10 seed borne fungi from rice seeds where Fusarium moniliforme, Curvularaia lunata, Aspergilus flavus and Rhizopus were most common encountered. Similar finding also reported by Butt et al. (2011) that the highest incidence of four fungal species namely, Fusarium moniliforme, Alternaria sp., Helminthosporium sp. and Curvularia sp. from different test rice varieties of Pakistan. Ora and co workers (2011) also reported that a total of 12 pathogens when blotter method, paper towel method and agar plate method were used to identify seed borne pathogens and among pathogens, Fusarium moniliforme Rhizopus stolonifer, Aspergillus sp. Bipolaris oryzae and Xanthomonas spp, were pre-dominant on all tested rice varieties from Bangladesh. Habib et al. (2012) reported that the highest percentage infection of Helminthosporium oryzae and Curvularia spp. from Pakistan as compared with other seed borne fungi when tested by agar plate and blotter paper method. A total of 69 rice seed samples from different states of India were tested their health status and sixteen genera of fungi viz. Acremonium, Alternaria, Aspergillus, Bipolaris, Chaetomium, Cladosporium, Curvularia, Exserohilum, Fusarium, Microdochium, Nigrospora, Phoma, Pyricularia, Rhizoctonia, Rhizopus and Verticillium comprising 27 species were found to be associated with the rice seed samples (Archana and Prakash 2013). Ahmed et al., (2013) detected many pre dominant fungi from 36 rice seed samples like Fusarium oxysporum, F. moniliforme, Bipolaris oryzae, Alternaria padwickii, Curvularia lunata, Aspergillus flavus, Aspergillus niger, Penicillium sp. and Nigrospora oryzae. In another set of experiment, agar plate method, 18 and 13 seed borne mycoflora were identified associated with farmer saved hill rice seed and improved varieties, respectively. These were Acremoniella sp. Alternaria padwicki, Chaetomium sp., Cladosporium sp., Curvularia lunata, C. oryzae, Dreschslera Fusarium moniliforme, oryzae, F. Pyricularia grisea, Rhizoctonia oxvsporum. sp., Saracladium oryzae, Tilletia sp., Trichothesiums sp., Aspergillus flavus, A. niger, Penicillium sp. and Rizopus stolonifer (Table 3). The highest incidence of Fusarium moniliforme was observed at range of 54.0-82.0% on all farmer saved hill rice seed and the maximum incidence recorded on Vai Buh (82%) whereas, the least incidence on RCM-10 (15.00%) a improved variety. Maipum (3) and Idaw (14) were recorded the least and highest number of pathogens genera, respectively.

Fusarium moniliforme was only the pre dominant seed borne pathogen recorded in all rice seed samples. Earlier workers have also reported various seed borne pathogens, Alternaria padwickii, Curvularia oryzae, C. lunata, B. oryzae, Aspergillus niger, Fusarium moniliforme, F. semitectum, F. solani and species of Phoma, Cercospora, Chaetomium, Sclerotium, Penicillium and Myrothecium from seeds of different varieties of rice in many parts of world (Wahid et al. 2001; Khan et al. 2001; Javaid et al. 2002; Nguefack et al. 2007; Utobo et al. 2011). Agarwal et al. (1989) also reported that fungi associated with Curvularia and Fusarium species which are known to cause leaf spot, pecky rice (kernel spotting) and root rot diseases in rice. Islam and Borthakur (2012) analyzed Aijung, rice variety of Assam, India for detection seed borne fungi by blotter method and agar plate method showed that species of Aspergillus, Fusarium, Alternaria and Curvularia are the dominant. In case of rolled paper towel method, the highest seed germination (92.26%) was observed on Maipum and the lowest seed germination (45.45%) on Idaw (Table 4). Maipum and Manipur Nem showed the highest vigour index of 1314.40 and 1172.27, respectively, whereas lowest vigour index was recorded on Idaw (37.42). These findings indicate that percent seed germination was decreased due to directly associated with seed borne pathogenic infection. The inferiors seed health, seed germination and seedling vigour of tribal farmers' saved hill rice seed may be improved through skill development of farmers on seed sorting and storage practices (Haque et al. 2007 and Kumar et al. 2013). Haque et al. (2012) reported that seedlings raised from cleaned seeds and their farmer saved seeds of the same variety results show a significantly higher grain yield in the cleaned seed than the farmer-saved seeds. Using poor quality rice seeds for planting reduces the productivity of landraces in attaining its genetic potential (Mew et al. 2004).

#### Conclusions

From the study it can be concluded that all rice seed of hill farmers collected from Mizoram, Northeastern of India carry a heavy load of seed borne of mycoflora which are responsible for loss in seed germination and seedling vigour except, *Maipum*. Since, rice is a staple food crop of this region; better seed health management and enhancement the seed replacement rate with quality seed is a prerequisite for successful rice cultivation by tribal peoples.

Pathogens							IIII fice see		,	0			nce seed		coflora									
				Jhun	1 Rice				Lowland (valley) rice									Improve cultivars						
	Idaw	Ak buh	Zakew	Saii Buh	Zaitlai	Vai buh	Zaizpui	Fazai	M (Rum)	Vui tawi	Buh tawi sang	Buh Mui	Shan buh	Sawkar Buh	Thlarau Buh	Zoro	Thinglang vai Buh	Tauphai Buh	M (Nem)	Maipum	RCM- 9	RCM- 10	RCM- 11	CAU R-1
Acremoniella	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	1.25	0.25	1.00	0.00	0.00	2.25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Alternaria padwicki	1.50	1.00	2.50	0.25	2.50	1.30	0.25	0.18	0.25	1.26	1.35	3.21	3.50	2.00	0.25	0.75	1.25	1.45	3.10	0.00	0.15	0.00	0.00	0.00
Chaetomium sp.	0.25	1.90	1.01	0.40	2.30	1.50	0.25	1.75	2.50	0.00	0.00	0.00	0.00	2.30	1.25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Cladosporium Curvularia lunata	0.00 2.00	0.00 4.00	0.00	0.00 5.25	0.00 6.25	0.00 4.00	0.00 5.50	0.00 3.00	0.00 2.00	0.00	0.00 7.25	0.00 3.00	2.10 2.00	5.15 7.50	0.00 3.00	0.00	0.00 2.25	5.25 5.50	3.25 3.20	0.00	0.00	0.0	0.00	0.00
Curvularia oryzae	0.25	0.50	0.00	0.00	0.00	0.00	0.00	0.00	0.25	2.00	0.50	3.00	0.25	3.00	2.00	1.00	1.50	1.00	2.15	0.00	0.75	7.15	7.50	1.00
Dreschslera oryzae	2.00	3.10	2.20	3.40	2.10	1.50	0.75	1.25	1.88	1.25	0.00	3.10	2.20	1.50	1.60	1.80	1.90	2.10	3.10	0.00	0.00	0.00	0.00	0.00
Fusarium moniforme	70.0	65.0	75.0	70.0	71.0	82.0	75.0	76.0	87.0	75.0	54.0	70.0	71.50	72.0	71.0	75.0	74.0	72.00	65.0	20.5	32.0	15.0	31.0	24.0
Fusarium oxysporum	3.50	3.10	1.50	1.30	3.50	4.50	0.25	0.10	0.25	0.30	0.40	2.50	0.00	0.00	3.10	3.00	1.75	3.00	1.50	0.23	3.30	1.75	1.25	1.00
Pyricularia grisea	2.00	1.50	1.75	0.25	1.75	1.00	0.25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	1.25	1.75	0.00	2.25	0.00	0.00	0.00	0.00	3.50
Rhizoctonia Saracladium oryzae	1.01 0.00	1.80 0.00	5.60 0.00	2.30 0.00	1.30 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.50	0.25	1.25 0.25	1.30 1.50	0.00	0.00 3.75	0.00	0.00	0.00	0.00	0.00
Tilletia sp. Trichothesium sp.	0.00 7.50	0.00 7.33	0.00 5.40	0.00	0.00 0.23	0.00	0.00	0.00	0.00	0.00 3.10	0.00 2.70	0.00 3.40	0.00	0.25	0.90 4.50	1.50 1.50	0.00 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Aspergillus flavus	5.50	4.20	6.10	3.00	3.40	2.00	3.10	1.00	1.20	3.00	0.00	3.20	1.50	0.75	0.15	2.00	3.00	1.00	2.10	0.00	3.10	2.10	3.10	2.10
Aspergillus niger	1.25	0.25	1.50	0.00	0.25	0.00	0.00	0.00	0.00	0.00	1.25	0.00	0.00	0.00	0.00	0.25	1.50	0.75	0.25	0.00	1.75	0.25	0.21	0.00
Penicillium sp. Rhizopus stolonifer	2.15 2.10	0.00 3.90	1.50 2.00	0.00	0.00 3.10	3.00 3.40	3.25 3.60	2.75 3.25	3.25 3.50	0.00	0.00 3.30	0.00 3.50	0.75 3.10	1.50 2.50	1.25 0.75	0.75 4.25	1.25 3.15	0.45	0.70 3.10	0.00	0.50 3.12	0.15	2.10 2.15	0.15
Nos .of Genera species	14	13	13	11	12	11	10	9	11	9	8	9	9	13	13	15	13	11	13	3	9	8	8	8

Table 3. Incidence of different seed borne mycoflora of farmer-saved hill rice seed of Mizoram, NE Region of India by Agar Plate Method

M=Manipur, RCM=Research Centre Maniphou, CAU-R=Central Agricultural University Rice

Name of Local races / Improved	Seed germination	Shoot length (cm)	Root length (cm)	Vigour index
Variety	(%)			
Idaw	45.45m	0.66k	0.17k	37.42j
Akbuh	76.99c-h	2.59fg	1.52hijk	316.17fghi
Zakew	71.24f-i	1.10jk	1.17hijk	161.25ij
Manipur (Rum)	86.78abc	4.84bc	6.56bc	988.96b
Vuitawi	75.03e-h	3.19ef	1.79hijk	373.38fg
Buh tawi sang	66.31hij	4.79c	5.16cde	659.77c
Buh Mui	80.10bc-h	2.87fg	2.01ghijk	390.35fg
Shan buh	66.75ej	3.89de	1.33hijk	348.67fgh
Saii Buh	75.43d-h	4.43cd	3.83fg	622.80cd
Sawkar Buh	79.14b-f	2.80fg	2.82fghi	444.26def
Zaitlai	50.10lm	4.11cd	1.69hijk	290.25fghi
Vai buh	80.00bcdef	2.17ghi	3.08fgi	419.73efg
Zai pui	72.84fghi	1.75hij	0.05ijk	192.05hij
Fazai	73.70efghi	2.42fgh	0.88ijk	390.35ghi
Thlarau Buh	89.36ab	3.20ef	3.78efg	623.11cd
Zoro	85.96abcd	2.17ghi	2.11ghij	367.61fgh
Thingtlang vai Buh	63.58ijk	2.39gh	6.19bc	545.49fgh
Tauphai Buh	77.20cdefg	1.56ij	0.48jk	157.49ij
Manipur (Nem)	84.60abcde	5.60b	8.25ab	1171.76a
Maipum	92.26a	7.41a	6.87a	1317.43a
RC Maniphou-9	80.57bcdef	2.14ghi	5.16cde	588.41cde
RC Maniphou-10	80.00bcdef	2.33ghi	5.16cd	599.20c
RC Maniphou-11	58.92ijkl	2.64fg	2.56def	306.38efg
CAU R-1	54.60klm	7.68a	4.65bc	673.22b
SEm+	5.3980	0.3936	0.9628	90.5094
CD ( $p = 0.05$ )	10.8539	0.7913	1.9359	181.9911

Table 4. Seed germination and seedling vigour of farmer-saved hill rice seed of Mizo	oram, NE Region of India by paper
towel method	

RCM=Research Centre Maniphou, CAU-R=Central Agricultural University Rice

Data presents the mean of three replications; Three hundred seeds were tested for each sample Figure with common letters did not differ significantly at 5% level by LSD.

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# References

- Abdul Baki AA, Anderson JD (1973). Vigour determination in soybean seed by multiple criteria. Crop Science 13: 630-633
- Agarwal PC, Mortensen CN, Mathur SB (1989). Seed borne diseases and seed health testing of rice. Phytopath Papers, p 106

Agrawal RL (1999). Seed Technology. 2<sup>nd</sup> Edn. Oxford and IBH Publishing Co. New Delhi, India, pp: 87-97.

- Ahmed M, Hossain M, Hassan K, Dash CK (2013). Efficacy of different plant extract on reducing seed borne infection and increasing germination of collected rice seed sample. Univ J Plant Sci 1: 66-73
- Archana B, Prakash HS (2013). Survey of seed-borne fungi associated with rice seeds in India. Int J Res Pure and Applied Microbiol 3: 25-29
- Atwal SS (2013). Successful entrepreneurship through seed production. In: Kumar A, Kumar R, Gupta A, Atwal SS (eds.) Entrepreneurship development through seed production, Indian Agriculture Research Institute, Regional Station, Karnal, Haryana, India pp:1.

- Barneet HL, Hunter BB (1972). Illustrated genera of Imperfect Fungi. 3<sup>rd</sup> Edn., Burgess Publishing Co., Minneapolis Minn, pp 241.
- Butt AR, Yaseen SI, Javaid D (2011). Seed-borne mycoflora of stored rice grains and its chemical control. J Animal Plant Sci 21: 93-196
- Habib A, Javed N, Sahi ST, Waheed M (2012). Detection of seed borne mycoflora of different coarse and fine rice varieties and their management through seed treatments, Pak J Phytopath 24: 133-136
- Haque A.H.M.M., Akon MAH, Islam MA, Khalequzzaman KM, Ali MA (2007). Study of seed health, germination and seedling vigour of farmer produced rice seeds. Int J Sustainable Crop Produc 2: 34-39
- Haque A.H.M.M, Elazegui FA, Taher Mia MA, Kamal MM, Haque MM (2012). Increase in rice yield through the use of quality seeds in Bangladesh. Afri J Agril Res 7: 3819-3827
- Ibiam OFA, Umemchuruba CI, Arnize AE (2006). Seed borne fungi associated with seed of rice (*Oryza* sativa L.) in storage and from the field in Ohaozara and Onicha local government areas of Ebonyi state. World J Biotech 7: 1062-1072
- Islam NF, Borthakur SK (2012). Screening of mycota associated with Aijung rice seed and their effects on seed germination and seedling vigour. Plant Patho Quarantine 2: 75–85
- ISTA (1999). International rules for seed testing. Seed Sci Tech 27: 333
- Janardhana GR, Raveesha KA, Shetty HS (1998). Modified atmosphere storage to prevent mould induced nutritional loss in maize. J Science Food Agri 76: 73-578
- Javaid MS, Wahid A, Idress M, Gill MA, Saleem A (2002). Seed mycoflora studies in rice. Pak J Phytopatho 14: 132-134
- Kavitha R, Umesha S, Shetty HS (2005). Dose dependent impact of dominant seed borne fungi on seed germination and seedling vigour of cotton seeds. Seed Res 33: 187-194
- Khan TZ, Gill MA, Khan MG (2000). Seed-borne fungi of rice from central Punjab and their control. Pak J Phytopathology 12: 12-14
- Kumar P, Singh PB, Gupta A, Maheshwari VK (2013). Safe seed storage: An essential component for successful seed entrepreneurship. In: Kumar A, Kumar R, Gupta A Atwal SS (eds.) Entrepreneurship development through seed production, Indian Agriculture Research Institute, Regional Station, Karnal, Haryana, India pp:28-29

- Mathur SB, Konsdal O (2003). Common laboratory seed health testing methods for detecting fungi. Bassersdorf, Switzerland, ISTA, 425pp
- Mew TW, Gonzales P (2003). A Handbook of Rice Seedborne Fungi. Laguna, Philippines/Enfield, NH, USA: International Rice Research Institute/Science Publishers, Inc. 83pp.
- Mew T., Leung H, Savary S, Vera Cruz C.M, Leach JE (2004). Looking ahead in rice disease research and management. Crit Rev Plant Sci 1: 1-25
- Nguefack J, Nguikwie SK, Fotio D (2007). Fungicidal potential of essential oils and fractions from *Cymbopogon* citrates, *Ocimum gratissimmum* and *Thymus vulgaris* to control *Alternaria padwickii* and *Bipolaris oryzae*, two seed-borne fungi of rice (*Oryza sativa* L.). J Essential Oil Res 17: 581-587
- Ou S.H (1985). Rice Diseases. CAB International Mycological, Institute Kew, Surrey, UK.395pp.
- Raj MH, Niranjana SR, Nayaka SC, and Shetty HS (2007). Health status of farmer's saved paddy, sorghum, sunflower and cowpea seeds in Karnataka, India. World J Agricultural Sci 3: 167-177
- SAS Institute Inc. (2011). SAS® 9.3 System Options: Reference, Second Edition. SAS Institute Inc., SAS Campus Drive, Cary, North Carolina.
- Sharma HL, Randhawa HS, Kapurm A, Singh S (1987). Seed discoloration in rice. Seed Research Production Unit 24: 37-41.
- Utobo EB, Ogbodo EN, A.C Nwogbaga (2011). Seedborne mycoflora associated with rice and their influence on growth at Abakaliki, Southeast Agro-Ecology, Nigeria. Libyan Agriculture Research Center Journal International 2: 79-84
- Wahid A, Javaid MS, Idress M, Gill MA (2001). Studies on the association and effect of *Fusarium solani* on rice seeds in Punjab, Pakistan. Proceeding 3rd National Conference of Plant Pathology, NARC Islamabad. pp. 70-72
- Warham EJ (1990). Effect of *Tilletia indica* infection on viability, germination and vigour of wheat seed. Plant Diseases 74: 130-135
- Zope AV, Thrimurty VS (2004). Effect of botanical pesticide on seed, rhizosphere microflora and seedling vigour in rice. J Mycol Plant Patho 34: 576–578