USE OF INDIGENOUS ISOLATES OF METARHIZIUM, ISARIA AND BEAUVIERA AS POTENTIAL BIO-CONTROL AGENTS AGAINST 
Sitophilus oryzae UNDER LABORATORY CONDITIONS

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Entomopathogenic fungi (EPF) are suggested as a new class of alternates followed by Synthetic chemical control of the insect pests. Virulence of seven EPF isolates, four strains of Metarhizium [(Qin-08, Qin-13, Qin-18 and ME-38 (LT-178)], two of Isaria {ME-33 (ILT-01), Yulin-5 (YL-01)} and one of Beauveria bassiana (Qin-21) were assessed against the adults of Sitophilus oryzae under laboratory conditions, firstly at single conidial concentration (1x10⁶ ml⁻¹) by immersion and the food mix methods. Qin-21, ME-33 and Qin-18 caused significantly highest mortality of S. oryzae at immersion methods (100, 100 and 98%), followed by food mix method to 100, 84.16 and 91.66% respectively. Yulin-5 was the least effective, showing significantly the lowest mortality at food mixed (32.49%) and immersion methods (40.20%) respectively. Hence, the immersion method was found to be most effective, resulting the higher mortality rate of S. oryzae in comparison to food mix method, in all tested fungal isolates. Secondly, we screened out the most effective isolates for multiple dose comparison i.e., 1x10⁶ to 1x10⁵ conidia ml⁻¹ by immersion method only. The isolate ME-33 resulted in 100% mortality of the pest at higher conidial dose as compared to Qin-21 and Qin-18 which showing 80 and 64.64% mortality respectively. The LT₅₀ was observed to be 3.63, 4.17 and 8.58 days in ME-33, Qin-21 and Qin-18, respectively at the highest conidial concentration (1x10⁷ ml⁻¹). ME-33 isolate with the highest mortality and lowest LT₅₀ at conidial concentration 1x10⁷ ml⁻¹ proved to be most effective for the control of S. oryzae. So these fungal isloates could be a better alternative for the management of S. oryzae. 

Keywords: Biological control, Entomopathogenic fungi, Pathogenicity, Sitophilus oryzae.

INTRODUCTION

Cereals are one of the essential sources of direct intake of calories for humans and also as major feed of animals. Cereals also afford a wide range of food components and nutrients like phyto-chemicals (Liu, 2007). Grains are attacked by a diversity of different insect pest’s species during their storage. The Post-harvest losses caused by these insects are estimated to be up to 20% or more in developing countries (Phillips and Throne, 2010; Rojht et al., 2010; Stejskal et al., 2015). The majority of the control measures against the storage pests mostly rely on application the synthetic residual chemical based insecticides, which are used at the time of their storage for long-term protection. In addition, various toxic fumigants, such as phosphine, are also applied as control measures (Hertlein et al., 2011). However, only a limited number of chemicals are commercially registered against the storage product pests (Yacoub and Nickolos, 2017).

Indiscriminate and un-judicial use of these types of chemicals have been restricted due to their toxicity, human health concerns, need to protect the environment, overall safety concerns, besides to their harmful influence on non-target organisms (Boyer et al., 2012; Yacoub, 2018). In addition, there is a problem of resistance development in insects to these different synthetic chemical insecticides which are recently being practiced (Butt et al., 2006). Fungi invading dead insects are called saprophagous, while fungi infecting the living insects are identified as entomopathogenic (Hibbett et al., 2007). Out of 1.5-5.1 million fungi species globally (Leger and Wang, 2010), Approximately 750-1000 fungi are entomopathogens, distributed in about 100 genera (Barra et al., 2013). Among various insect pests of stored commodities, the Rice weevil, Sitophilus oryzae (Coleoptera: Curculionidae) is considered a most notorious pest specie of the storage commodities worldwide (Kaur et al., 2014). The adults of S. oryzae feed on grains while larvae prefer to feed inside the grain kernels and are responsible for both the quantity and quality losses of rice grain in stores (Kavallieratos et al., 2014; Nguyen et al., 2016). Weevils Population has developed the resistance against many chemical pesticides as well as fumigants which is commonly applied against the storage pests (Butt et al., 2006; Ramanujam et al., 2014).

Recently, more than 700 species and 90 genera of fungi were observed for infecting insects (Batta, 2008; Ramanujam et al., 2015; Maina et al., 2018). Most of these genera belong to
family Deuteromycetes and Entomophthorales (Lord, 2007). *Beauveria bassiana* sensulato (Balsamo) Vuillemin (Ascomycota: Hypocreales) is used for the organization of stored products pests (Batta, 2005). *B. bassiana* has been reported as highly effective beside the major stored-product insect’s viz., *S. oryzae*, *R. dominica*, and *T. castaneum* (Kaur et al., 2014; Kavallieratos et al., 2014; Nguyen et al., 2016). The EPF application is relatively safe against the coleopteran pest for the protection of stored items. Use of EPF for the control of *S. oryzae* is very rare and only few studies were conducted with the fungi *Metarhizium anisopliae* and *Beauveria bassiana* for the management of *S. oryzae* (Sanchez-Rodriguez et al., 2017; Pedrini et al., 2007). EPFs penetrate the insect body via cuticle, causing insect mortality. Fungi then inside insect body germinate and recycle on the cadavers, thus returning more inoculums to the stored-product system (Tadele and Pringle, 2007; Zhang et al., 2008; Stephehu et al., 2012; Ortiz-Urquiza and Keyhani, 2013; Mandira et al., 2018). The present study indicated that selection of the virulent isolates was the most important in using entomopathogenic fungi for biological control of *S. oryzae*. The objective of present work was to investigate the pathogenicity of seven entomopathogenic fungi isolates such as *Metarhizium anisopliae* (Qin-08, Qin-13, Qin-18 and ME-38 (LT-178), two of *Isaria cateniannulata* ME-33 (ILT-01), Yulin-5 (IYL-01) and one *Beauveria bassiana* (Qin-21), against the adults of *S. oryzae* under laboratory conditions.

**MATERIALS AND METHODS**

**Insect culture:** The Preliminary Population of *S. oryzae* was collected from insect-infested rice store at Yangling, Shaanxi Province (China). The stock was maintained under the laboratory conditions at Insect Related Resources (LIRR), Northwest A & F University (NWAFU). The test insects were reared at 28°C with RH 65% (Tadesse and Subramanyam, 2018), in 2-liter glass jars having 1000-g rice grains free from any contamination. The transparent glass jars were covered with a piece of fine muslin’s cloth. Two weeks later, original adults from the jars were removed out. The jars were observed daily and the same aged newly emerged 30 adults were used each for all series of bioassays.

**Source of fungal isolates:** Entomopathogenic fungi isolates were provided by Laboratory of insect related resources (LIRR) College of Plant Protection, Northwest A&F University (Table 1).

**Preparation of fungal conidia:** For preparation the fungal suspensions of the seven fungal isolates, single spore isolates were cultured on Sabouraud-dextrose-yeast extract agar medium (SDAY) with 3.5-4 mm depth. The petri plates then kept in incubator for the period of two weeks, and the plates were without Para film lid sealing and by putting them in total darkness. They were put on 26°C and then shifted it in incubator with the relative humidity (RH) 70% for a week more with 14:10 light and dark period. Sterilized spatala was used for the conidal harvesting at once. Prior to use in sterile 0.02% Tween 20 (Tianjin Guangfu Fine Chemical Research Institute, Xi’an, China).and then vortexed for two minutes. To minimize the Tween influence, fungal suspensions stock solution was diluted with sterilize ddH2O. Concentrations of conidial suspensions were adjusted from 1x10⁴ to 1x10⁸ conidia ml⁻¹ by using an improved-Neubauer hemocytometer (Lauda-Königshofen, Germany) with a magnification which range 20X by the use of inverted microscope (Nikon Eclipse TE-2000-S Japan). Germination rate of conidia for all the tested isolates were assessed by planting the conidia 100 microliter in a 10⁷ conidia ml⁻¹ suspension with a quarter ratio of SDAY at 26 ±2 in a complete darkness (Inglis et al., 1997). **Initial pathogenicity assay for EPF isolates against *S. oryzae***: Firstly, the single highest concentration 1x10⁸ conidia

<table>
<thead>
<tr>
<th>Fungal species</th>
<th>Isolate code</th>
<th>Origin</th>
<th>Isolation host</th>
<th>Altitude</th>
<th>Year</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Metarhizium robertsii</em></td>
<td>Qin-08</td>
<td>FNNR, Qinling Mountains Southwest Shaanxi province, China</td>
<td>Soil selective medium</td>
<td>107°40′21″E, 33°34′12″N (1000 m)</td>
<td>2015</td>
</tr>
<tr>
<td><em>M. anisopliae</em></td>
<td>Qin-13</td>
<td>-do-</td>
<td>-do-</td>
<td>107° 47′ 14″ E, 33° 39′ 79″ N (2000 m)</td>
<td>2015</td>
</tr>
<tr>
<td><em>M. pingshaense</em></td>
<td>Qin-18</td>
<td>-do-</td>
<td>-do-</td>
<td>107° 55′ 13″ E, 33° 45′ 81″ N (3000 m)</td>
<td>2015</td>
</tr>
<tr>
<td><em>M. robertsii</em></td>
<td>ME-38 (LT-178)</td>
<td>Sichuan province, (Bazhong, Ling Ton) China</td>
<td>Bait insect method (mealworm larvae, <em>Tenebriotomolitor</em>)</td>
<td>30° 15′ 80″ N, 100° 58′ 67″ E (3380m)</td>
<td>2016</td>
</tr>
<tr>
<td><em>Isaria cateniannulatus</em></td>
<td>ME-33 (ILT-01)</td>
<td>Sichuan province, (Bazhong, Ling Ton) China</td>
<td>-do-</td>
<td>30° 15′ 80″ N, 100° 58′ 67″ E (3380 m)</td>
<td>2016</td>
</tr>
<tr>
<td><em>Beauveria bassiana</em></td>
<td>Qin-21</td>
<td>FNNR, Qinling Mountains Southwest Shaanxi province, China</td>
<td>-do-</td>
<td>33° 39′ 79″ N, 107° 47′ 14″ E (1600m)-</td>
<td>2015</td>
</tr>
</tbody>
</table>
ml⁻¹ was tested for pathogenicity (virulence). Two methods of fungus application were studied: 1- The immersion or dip method, and 2- An application with food mixture. In both methods, the test insects and food were immersed for one minute into the conidial suspension of 5-ml and the excess of conidial excessive suspension were removed by putting the test insects and a food on a sterilize filter paper (Dong yang, China). All the insects, treated in the independent biological repeats were later shifted to WD9-3.5-cm petri plates (Zhejiang Plasmid Medical Technology Co., Ltd., China). All petri plates were settled under the complete randomized design (CRD) and were kept under the Standard climatic chamber environment of 26 ± 2°C, 80 ± 7% RH with photo phase duration of 12-h:12-h (L: D). All plates were partially sealed with plastic solution tape to avoid the interference of insects. The control adults were treated with sterilized ddH₂O. Mortality data were recorded every 24 hours for 14 days.

**Virulence assay of EPF against S. oryzae**: Based on results of the above-mentioned initial Pathogenicity assay, the isolates that showed the highest mortality were selected for further evaluation of their virulence against the S. oryzae. Four different concentrations of conidia were used for each isolate i.e., 1x10⁴, 1x10⁵, 1x10⁶ and 1x10⁷ ml⁻¹. Sterilized ddH₂O aqueous solution was used for control. For the virulence test, an immersion method was used following the same procedures, 5-ml conidial suspensions. Fifteen Sitophilus oryzae adults were used for every replication and the mortality data were recorded for 24-hours each up to 15-days. Mean survival period for each isolate at each concentration was also determined.

**Statistical analysis**: All the experimental assays in this study were performed by using thrice biological repeats of each fungal isolate under Complete randomized design (CRD). Mortality data were recorded. An (ANOVA) Analysis of variance was used to examine the significant differences statistically within the tested isolates. Obtained data was analyzed statistically by using SAS 13.2. Probit analysis was used to calculate the mean survival times (MST). All required graphs were prepared by using excel software.

**RESULTS**

**Fungal Isolates tested by Immersion Method at Single Concentration Bioassay**: The results presented in (Fig.1) showed that all the isolates differed significantly (P≤0.03). The mortality of the test insect was ranged from 40.2 to 100 percent. The treatments ME-33 and Beauveria bassiana (Qin-21) showed the highest mortality (100%) and did not show significant difference with Qin-18 (91.66%). Qin-18 also showed non-significant variation with those of ME-38 (88.22%) and Qin-13 (86.66%). The later mentioned treatments were also at par statistically with Qin-8 (68.5%) and found to be intermediate. The lowest mortality was observed in Yulin-5 (40.2%) and differed significantly from all the treatments. The treatments Beauveria bassina (Qin-21), ME-33 and Qin-18 proved to be the most effective while Yulin-5, the least.

![Figure 1. Mortality % of S. oryzae adults tested to immersion method at 1x10⁶ conidia ml⁻¹.](image)

**Fungal Isolates tested via Food Mix Method at Single Concentration Bioassay**: Significant difference (P≤ 0.002) was found to exist among treatments (Fig.2). Beauveria bassiana (Qin-21) showed the highest mortality (100%) of S. oryzae and was at par statistically with those of observed in ME-33 and Qin-18 showing pest mortality 98.00 and 84.16%, respectively. The lowest mortality (32.49%) was found in Yulin-5 and did not show significant variation with those of found in ME-38 (33.33%), Qin-8 (51.8%) and Qin-13 (54.1%). The treatment Qin-18 was also at par statistically with Qin-8 and Qin-13. Conclusively, Beauveria bassiana (Qin-21) ME-33 and Qin-18 were the most effective while Yulin-5 and Qin-8 the least against S. oryzae. Qin-13 and Qin-8 were classified as intermediate.

![Figure 2. Mortality (%) of S. oryzae adults exposed to food mix method at 1x10⁶ conidia ml⁻¹.](image)

**Comparison of Immersion and Food Mix Fungal Isolates**: The results (Fig. 3) revealed that significantly (P ≤ 0.001) maximum mortality (100%) of the pest was found in
*Beauveria* at both application methods and was at par statistically with those of observed in ME-33 (100%) at immersion method. The lowest mortality of the pest was observed in Yulin-5 at both methods compared with the remaining pathogens. Immersion method showed higher mortality of the pest in all the isolates as compared to food mixed method.

**Multiple concentration bioassays**: As per initial screening assays, most effective fungal isolates against *S. oryzae* screened out from single concentration bioassay test were studied for multiple concentration bioassays by immersion method. The results revealed significant variation (*P*<0.001) among different concentrations in all the fungal isolates (Fig. 4, 5 and 6). Higher conidial concentrations resulted in maximum mortality of the pest as compared to lower concentrations in all the fungal isolates. The observed mortality of the pest ranged from 44.44 to 80.00, 35.56 to 100 and 40.00 to 64.64 percent at conidial concentration 1x10⁷ to 1x10⁴ in *Beauveria bassiana* (Qin-21), ME-33 and Qin-18, respectively. The results presented in Table 1 revealed that ME-33 showed less LT₅₀ i.e., 3.63 days with a range of 3.41 to 3.83 days at higher conidial concentration. The comparison of different isolates (Fig.7 and Table 2) revealed that ME-33 was found to be the most reliable fungal isolate to use against *S. oryzae* at 1x10⁷ conidial concentration compared with the other isolates. All the isolates; however had a potential to control the pest even at lower conidial concentrations by immersion method.

![Figure 3](image3.png)

**Figure 3.** Mortality (%) of *S. oryzae* adults tested to immersion and food mix method at conidial concentration of 1x10⁵.

![Figure 4](image4.png)

**Figure 4.** Mortality (%) of *S. oryzae* adults tested to *Beauveria bassiana* at different concentrations.

![Figure 5](image5.png)

**Figure 5.** Mortality (%) of *S.oryzae* adults tested to ME-33 at different conidial concentrations.

<table>
<thead>
<tr>
<th>Conidial Conc. (ml⁻¹)</th>
<th>(Qin-21) Percentile</th>
<th>95% Fiducial CI</th>
<th>(ME-33) Percentile</th>
<th>95% Fiducial CI</th>
<th>(Qin-18) Percentile</th>
<th>95% Fiducial CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 x 10⁷</td>
<td>4.17±0.18</td>
<td>3.79</td>
<td>4.53</td>
<td>3.63±0.11</td>
<td>3.41</td>
<td>3.83</td>
</tr>
<tr>
<td>1 x 10⁶</td>
<td>11.52±0.64</td>
<td>10.40</td>
<td>13.01</td>
<td>6.40±0.21</td>
<td>5.98</td>
<td>6.82</td>
</tr>
<tr>
<td>1 x 10⁵</td>
<td>17.82±1.65</td>
<td>15.18</td>
<td>22.05</td>
<td>16.87±1.47</td>
<td>14.50</td>
<td>20.61</td>
</tr>
<tr>
<td>1x10⁴</td>
<td>19.78±1.92</td>
<td>16.75</td>
<td>24.78</td>
<td>20.32±2.10</td>
<td>17.05</td>
<td>25.90</td>
</tr>
</tbody>
</table>

Table 2. Comparative LT₅₀ at various conidial concentrations in selected virulent EPF isolates.
Potential use of Indigenous Entomopathogenic fungi against S. oryzae (L.)

**LT₅₀ at various conidial concentrations:** The results presented in Table 2 reveals that ME-33 at conidial dose 1x10⁷ was the most effective showing minimum LT₅₀ i.e. 3.63 days. The fiducial CI at 95% ranged from 3.41 to 3.83. The LT₅₀ values increased on decreasing the conidial concentration in all the isolates. The application of Qin-18 resulted in minimum effect on the pest as it inclined longer period to the pest for its survival with fiducial CI range from 7.96 to 9.30 at the highest conidial concentration. The virulent ME-33 at highest conidial concentration performed the best in controlling the pest as compared to (Qin-21) and (Qin-18). Thus ME-33 is recommended at conidial dose of 1x 10⁷ ml⁻¹ for the effective control of the pest under study and suggested to include this in IPM tactics.

**DISCUSSION**

In-vitro, studies were carried out to determine the comparative virulence of seven entomopathogenic isolates against S. oryzae on stored rice at single conidial concentration i.e., 1x10⁸ ml⁻¹ by food mixed and immersion methods. Beauveria bassiana (Qin-21) isolates, (ME-33) and (Qin-18) isolates were found to be the most effective and based on the high insect mortality and low MST values among both methods of application, isolates were selected for further studies at multiple conidial concentrations i.e., 1x10⁴ to 1x10⁷ by immersion method. The results of this study demonstrated significant variation among the different fungal isolates and genera. S. oryzae adults were susceptible to EPF isolates that

![Figure 6. Mortality (%) of S. oryzae adults exposed to Qin-18 at different conidial concentrations.](image)

![Figure 7. Mortality (%) of S. oryzae adults exposed to three different fungal isolates at different conidial concentrations.](image)

![Figure 8. Fungal mycosis symptoms on the beetle cadavers of three different genera of EPF, isolates, Isaria cateniannulatus ME-33 (ILT-01), B. bassiana (Qin-21) and M. pingshaense (Qin-18). Mycosis photos were taken by advance stereomicroscope system (Discovery V20, Zeiss; CCD, Axio, ICC5, ZEISS; and Leica 205C, Germany, (a-c); Conidiospores photos of the highly virulent isolates were taken by the SOPTOP-EX20, Biological microscope, Shanghai Jonchang Co., Ltd (d-f).](image)
were tested in this study. Toxicity of the isolates and its possible pathogenicity level depend on the bioassay conditions as well as the host immune system. In many of these type of studies, effective results were observed when stored grains were treated with entomopathogenic fungi, particularly *B. bassiana* and *M. anisopliae*. Similarly, this study also demonstrates significant results for the pathogenic isolates, which were very effective and responsible for *S. oryzae* mortalities (Athanassiou et al., 2007, 2008; Hansen et al., 2007; Batta, 2013), but they have different virulence levels. Furthermore, our results indicate difference in susceptibility among different fungi against *S. oryzae*. ME-33, Qin-21 and Qin-18 were found to be more effective among all tested fungi against *S. oryzae*. (Batta, 2016a;b; Batta, 2012), described the high degree of susceptibility of two different insects against *B. bassiana* and *S. maize* and found similar results in different fungus against larvae of *Plutella xylostella*. Consequently, our results are in agreement with earlier study by (Dal et al., 2006), exposed that differences in a fungal ability to reduce the host defense mechanism lead to treatments with a higher LT50 as well prolong the insects’ survival. The variation observed in the isolates, LT50 values likely reflects the physiological and genetic differences among these fungal isolates. 

The present study revealed that, conidial concentration was the primary factor which keeping up the “speed of kill” of the subjected insects in majority of observations. Despite the fact that there was no prior study available that compared these seven fungi against *S. oryzae*. Previous reports suggested that *B. bassiana* and *M. anisopliae* might be more effective than other isolates. Indeed, (Talaei et al., 2006; Batta et al., 2010; Batta, 2016a) reported that there is no reliable indicator for the virulence that shows the original host is specific for the specific fungus. Ramanujam et al. (2015) reported that the isolates PPRCH-HH and PPRC-4 showed the high pathogenicity against the stem borer *Chilo partellus*. In some other studies it has been reported that most virulent isolates are isolated from closely related host or the same place (Batta and Abu, 2005). It is also agreed that some fungus has broad host range like *B. bassiana* and *M. anisopliae* and their pathogenicity also varies as per their host. Subsequently, (Batta, 2007; Addis and Tadele, 2009) reported an increasing trend in *M. anisopliae* and *B. bassiana* that pathogenicity drives through the habitat selection. Therefore, the current results revealed that the selection of the potential isolates would not be restricted to isolates of fungi from the sole host mass. Entomopathogenic fungi isolates, ME-33 (ILT-01) (*Isaria cateniformis*) was consistently more virulent than *Beauveria bassiana* (Qin-21) and (Qin-18) *M. pingshaense*. Thus, considering the low mean survival time MST range, these fungi isolates could be reflected more pathogenic amongst all tested isolates against the rice weevil management Program. Maximum mortality rate was detected on the high conidial concentrations of 1×10^6 and 1×10^7 conidia/ml which shows that more number of conidia with the high concentrations as well as adhere chances are increased. Mortality was observed at all the concentrations of each isolate but this mortality is limited which ranged from 35.5% to 40% at lower concentrations and this low mortality is responsible for slow development of disease or infection which was the effect of low dose application (Batta, 2012). In the present study, high conidial concentration caused the mortality of insect in a short time (3.41 to 8.58 days) whereas, low conidial dose took a longer period (19.24 to 20.32 days). These findings confirm that the results of (Batta et al., 2010), who stated that the concentrations applied with high conidia rates caused the mortality in a short time but on the other hand doses with low conidial concentrations may take two or more weeks to kill the test organisms. Different *Beauveria bassiana* isolates Pathogenicity was evaluated against the corn weevil by (Addis and Tadele, 2009; Cherry et al., 2005) reported that among these ten isolates one isolates showed higher mortality (88%) within 8-days at the low concentration of 1×10^7 conidia ml^{-1}. In some other studies high dose requirement is reported against *T. Castenenum* and *Callosobruchus maculatus* (Clare et al., 2016). However, in some other studies low concentration doses against Stored Grain Beetles were required (Mehmet et al., 2016). George et al. (2018) examined that *B. bassiana* isolates were much lethal against *S. zeamais* only at high concentrations of 1×10^7 conidia ml^{-1} and reported that pathogenicity varies within the isolates was superficial. The method of application as well as fungal formulation was the most important factor in its efficacy against *S. oryzae* adults. Previous studies provide interesting information on this issue. For example, Cherry et al. (2005) reported 100% mortality of *C. maculatus* adults dipped for 5seconds into a conidial suspension of *M. anisopliae* and *B. bassiana* after 6 and 8 days of introduction. Furthermore, it has been reported by (George et al., 2018), stated *S. zeamais* adults which were dipped into certain isolates of *B. bassiana* suspension for 5 seconds suffer 100% mortality after only 4 days after application time. Nearly similar results were also found for *S. oryzae* treated with certain *B. bassiana* and *M. anisopliae* isolates. As for the application methods examined here, direct application (Immersion method) of the fungus on insect bodies is much successful compare to contact of insects with the fungi isolates which is applied on the food substrate.

**Conclusion:** The present study is novel tactic linking the use of indigenous isolates. Mortality results were found to be directly related to the conidial concentrations as well as the exposure time. Among the selected isolates, the highest mortality of 100, 80 and 64.64 % was observed on the conidial dose of 1×10^7 for the isolates ME-33, Qin-21 and the Qin-18. Tested isolates could be suggested as promising candidates in terms of virulence and pathogenicity for the integrated pest management program against *S. oryzae* and the local pest species of the stored commodities. Further research needed to
develop the myco-insecticides for the control of different pests. Considering different aspects like mass productions, storage, formulations, spectrum of activities against the non-target organisms under the various environmental conditions.

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