



## EFFICACY OF CARBENDAZIM AGAINST CHARCOAL ROT OF MAIZE

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### KEYWORDS :

#### Introduction:-

Maize (*Zea mays* L.) is a large grain plant domesticated by indigenous people in Mesoamerica in prehistoric time. It is best grown in warm, tropical regions as it requires warm soil to develop optimally. It is an annual grass belonging to Poaceae and is a staple food crop grown all over the world. It is also commonly grown as a feed for livestock.

Such an important crop suffers from many fungal diseases, such as common smut caused by *Ustilago maydis* (de Candolle) Corda, head smut caused by *Sphacelotheca reiliana* (Kunhn) Clinton, brown rot caused by *Physoderma zae maydis* F. J. Shaw, rust caused by *Puccinia sorghi* Schw., leaf blight caused by *Exserohilum turcicum* (Pass.) Leonard et Suggs. Seedling blight and wilt caused by *Fusarium moniliforme* var. *subglutinans* Wollenw and Reinking, seedling blight and top rot caused by *Giberella zae* (Schwein) Petch, Charcoal rot caused by *Macrophomina phaseolina* (Tassi.) Goid. Among these charcoal rot caused by *Macrophomina phaseolina* (Tassi.) Goid. is serious.

#### HEALTHY PLANTS OF MAIZE



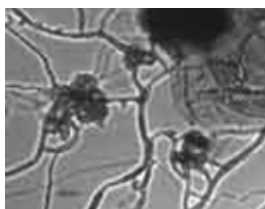
#### INFECTED PLANTS



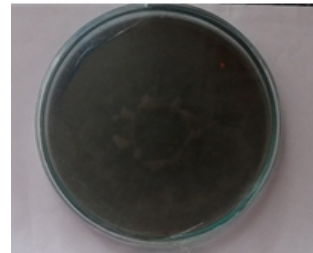
#### INFECTED STEM



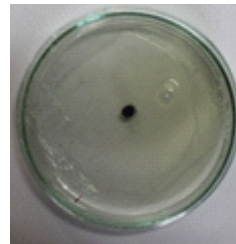
#### SCLEROTIA



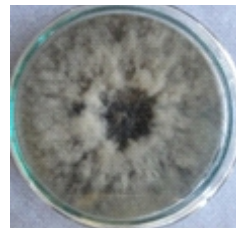
#### PURE CULTURE



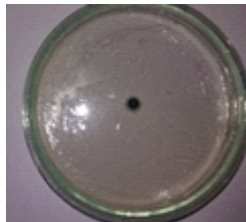
#### INVITROMIC SENSITIVE MP<sub>3</sub>



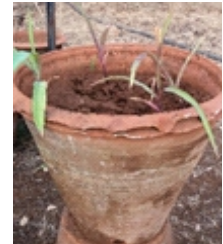
#### CONTROL



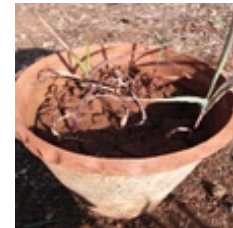
#### RESISTANT MP<sub>3</sub>



#### INVIVOMIC SENSITIVE MP<sub>3</sub>



#### CONTROL



#### RESISTANT MP<sub>3</sub>



#### Material and Method:-

16 samples exhibiting charcoal rot of maize were collected from different districts of Maharashtra viz. Kolhapur (Minche savarde), Sangli, Satara (Nagthane, Songaon, Sangavi) Pune (Baramati), Ahamadnagar (Ane), Solapur (Madha), Nasik (Dindori), Belgaon (Padalihal), Nanded (Bomnali) Latur (Chakur), Parbhani (Bramhanpur), Usmanabad (Umarga) Jalana (Dawalwadi) and Bid (Nirgudi). To isolate the causal agent, the collected samples were brought to the laboratory in sterilized bags. The infected portion of

stem is cut in to the size 2 mm and sterilized by using 0.1% HgCl<sub>2</sub>, and washed with sterilized distilled water (Jadhav et al., 2010), these sterilized stem portion were kept on a czapek dox agar plates amended with streptomycin sulphate (Patil et al., 2012; Mali et al., 2015). Inoculated plates were incubated at 30 ± 2° C for growth of the fungus and further studies (Mali et al., 2016). After 5-6 days of culture, black fungal mass was observed. On the basis of morphological, microscopic characters and following relevant mycological literature the fungal isolate was identified as *Macrophomina phaseolina* (Tassi.) Goid. In this manner, 16 isolates were obtained.

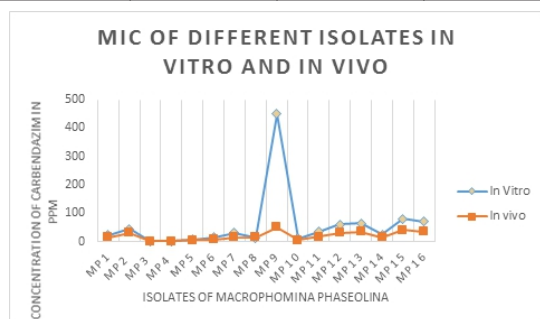
The *in vitro* sensitivity of *Macrophomina phaseolina* (Tassi.) Goid. was carried out by using Food Poisoning Technique (Dekker and Gielink, 1979). Czapek Dox agar medium plates were prepared containing different concentrations of carbendazim..

After solidification of media, a disc (6 mm) with fungal culture was obtained from the margin of an actively growing colony and placed upside down on the agar surface. These plates were then incubated at 30 ± 2° in 12 hour cycle of dark and light and then continuous growth was measured after various time intervals. Plates without carbendazim were served as control.

For *in vivo* experiments Mycelial suspensions of all fungal isolate were prepared in sterile distilled water, and then inoculated on the healthy plants of maize. treated with 10 ml solution of different concentrations of carbendazim 24 hours before the inoculation. The experiment was carried out in triplicates. The plants without fungicide treatment served as control. The plants without any treatment served as absolute control.

**Table 1:-** MIC (Minimum Inhibitory Concentration) of carbendazim against *Macrophomina phaseolina* (Tassi.) Goid. isolates causing charcoal rot of maize.

Locality	Isolate	<i>in vitro</i> (ppm)	<i>in vivo</i> (ppm)
Minche Savarde	MP <sub>1</sub>	22	15
Nagathane	MP <sub>2</sub>	44	30
Songaon	MP <sub>3</sub>	1	1
Sangavi	MP <sub>4</sub>	2	1.5
Ane	MP <sub>5</sub>	8	6
Madha	MP <sub>6</sub>	15	8
Dindori	MP <sub>7</sub>	30	14
Baramati	MP <sub>8</sub>	12	15
Padalihal	MP <sub>9</sub>	450	50
Sangali	MP <sub>10</sub>	10	6
Chakur	MP <sub>11</sub>	35	16
Bomnali	MP <sub>12</sub>	60	30
Bramhanpurpur	MP <sub>13</sub>	65	35
Umarga	MP <sub>14</sub>	25	15
Dawalwadi	MP <sub>15</sub>	80	40
Nirgudi	MP <sub>16</sub>	70	35



### Result and Discussion:-

There was variation in the minimum inhibitory concentration of carbendazim. MIC on agar plates ranged from 1 to 450 ppm and it was 1 to 50 ppm on maize plants. Isolate MP<sub>3</sub> was sensitive to carbendazim and it showed 1 ppm MIC both *in vitro* and *in vivo* While isolate MP<sub>9</sub> was resistant to carbendazim and showed 450 ppm MIC on agar plates and 50 ppm on maize plants. The results are in agreement with other workers also. Wadikar, et al (2008) There was large variation in the MIC of carbendazim among the *Macrophomina phaseolina* isolates both on agar plate and pigeon pea plant, MIC on agar plate ranged from 80 to 140 mg/ml, while it was 65-127 mg/ml on pigeon pea plant.

Similarly, Bhale (2009) reported the MIC of carbendazim against *Alternaria alternata* causing leaf spot of spinach was ranging from 350 to 700 µg/ml. According to Khandare (2013) the MIC of carbendazim among 12 isolates of *Alternaria alternata* causing root rot of fenugreek was ranging from 2500 to 5000 µg/ml *in vitro* and 500 to 1000 µg/ml *in vivo*.

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