

RESEARCH ARTICLE

Estimation of deteriorative effect of *Fusarium oxysporum* and *Aspergillus niger* on fenugreek seed germination, seedling vigour and *in vitro* efficacy of fungicides

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ABSTRACT

In the present investigation, seed and soil inoculations were used to prove the pathogenicity of *Fusarium oxysporum* and *Aspergillus niger*, isolated from fenugreek seeds. Out of both, *Fusarium oxysporum* proved to be highly virulent as it caused higher per cent pre and post-emergence mortality (14.50% and 8.75%) and seedlings showed 49.55 per cent symptoms, whereas *Aspergillus niger* was observed to be less pathogenic because it caused lower per cent pre and post-emergence mortality (6% and 4.55%) and seedlings showed (31.25%) symptoms. In *in vitro* test, complete inhibition was observed on Bavistin at all concentrations used. Whereas, 90.40 per cent growth inhibition occurred on Thiram at lowest concentration used i.e. 50 ppm. Captan and Raxil were not so effective at lower concentrations.

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INTRODUCTION

Fenugreek (*Trigonella foenum-graecum*), an annual legume native to the Mediterranean region, locally known as *Methi*, is cultivated not only as a leafy vegetable but also for medicinal purposes (Som and Maity, 1993). It is cultivated in counties India, Argentina, Egypt, Southern France, Morocco and Lebanon. Green methi is a good source of iron (Fe) as well as other minerals for human beings (Chhibba *et al.*, 2000). Seeds contain proteins 26 per cent, water-soluble polysaccharide (galactmannan) 20 per cent, hemi-cellulose and cellulose 24.5 per cent, water 9 per cent, fat (fenugreek oil) 7 per cent, lignin 2.5 per cent and saponin 8-10 per cent. The crop suffers severely from few seed-borne diseases including wilt caused by *Fusarium oxysporum* Schlecht which also affects seed germination, vigour index, plant growth and the grain yield (Shivpuri and Bansal, 1987 and Hashmi, 1988).

MATERIALS AND METHODS

Two tests were performed :

Soil inoculation technique :

Fusarium oxysporum and *Aspergillus niger* observed on fenugreek seeds were grown, separately on autoclaved rice medium (20g rice + 10 ml distilled water) contained in 250 ml conical flasks. These flasks were inoculated with spore/mycelial suspension prepared from 7 days old fungal culture. Flasks were shaken every day to avoid clumping. Autoclaved soil (Soil : FYM = 3:1, autoclaved at 1.045 kg/cm² for 1 hour for 3 consecutive days) was filled up in 30 cm earthen pots (pre-sterilized with 0.1 per cent HgCl₂ solution for 3 minutes followed by 3 washing with sterilized distilled water) and were inoculated with *Fusarium* sp. separately. For inoculation, the upper 4 cm layer of the soil was thoroughly mixed with rice medium supporting fungal growth. The pots were covered with polythene bags and left for 24 hours in a cage house