

Biological Control of Parthenium Weed

Abstract

A total of 12 fungi associated with the seeds of Parthenium were recovered. These includes severe highly plant pathogenic fungi viz. Colletotrichum, dematium, fusarium species, phoma herbarium, Drechslera sp, Curvularia lunata, Rhizopus herbarium and Colletotrichum- dematum were also responsible for severe seedling blight in Parthenium.

Some fungi associated with externally are well known for their capacity to produce phytotoxic secondary metabolites. Except few exceptions all of them significantly reduced the seed germination.

Keywords: Weed/Biological Control/Parthenium

Introduction

Parthenium (*P. hysterophorus* L.) is now considered as a serious weed in India and now considered as a serious weed in India and the problems due to the weed is very well documented by D.R. Rajesh Garg and D.R. Rajesh Singh. Seeds are known to carry various types of pathogenic as well as non pathogenic fungi, some are responsible for severe seed and seedling diseases, besides synthesizing potentially toxic secondary metabolites. Therefore, the present communication deals with the incidence of seed mycoflora and its impacts on seed germination.

Material and Methods

Fungi were isolated from infected seeds of *Parthenium* through Blotter method (ISTA3) and Dilution method. Pathogenicity test was carried out by D.R. Rajesh Garg and D.R. Rajesh Singh. The percent incidence of fungi and percent inhibition in seed germination were determined by using the following formula= $\frac{\text{Total no of seeds colonized by individual fungus}}{\text{Total no of seeds observed}} \times 100$
percent inhibition in seed germination = $\frac{\text{Germination in infected seed}}{\text{Germination in controlled seed}} \times 100$

All Experiments were Conducted in Triplicate

Result and Discussion

Twelve fungi were isolated from infested/infected seeds of PARTHENIUM (TABLE-1). *Aspergillus Fumigatus* (81.36%) was most dominant isolate which was followed by *A. niger*, *A. flavus*, *Fusarium rigidusculum*, *F. oxysporum* and *Drechslera biseptata*. Fungal isolates recovered from significant external surface and caused reduction in germinability of seeds. Significant reduction in viability and germinability due to externally associated fungi have also been recorded by many workers. This might be because of invasion of outer cover or secretion of toxic metabolites. Similarly some well known plant pathogenic fungi viz, *Fusarium rigidusculum*, *F. oxysporium*, *Phoma herbarium*, *P. hyunmicola*, *Drechslera biseptata*, *Curvularia luata* and *Colletotrichum dematim* were also found to be associated internally. Maximum inhibition in seed germination was due to *C. dematium* FGCC#20 (78.34%). It was followed by *Phoma herborum* (69.94%). Seed borne nature of these fungi also been recorded by some earlier workers.

On the basis of above observations it can be concluded that the seed mycoflora recorded have enormous mycoherbicidal potential against the hazardous weed Parthenium. Further investigation are to be carried out for their large scale exploitation.

R.K.Garg

Assistant Professor,
Deptt. of Botany,
Govt. P.G. College Satna,
M.P.

Rajesh Singh

Assistant Professor,
Deptt. of Botany,
P.G. College Maihar,
M.P.

| S.No. | % incidence of fungi | % inhibition of seed germination | | |
|----------------|-------------------------|----------------------------------|-------|-------|
| Blotter Melleo | | Dilution in the | | |
| 1 | Apergillus Niger | 0.0 | 79.28 | 22.00 |
| 2 | A.Fumigatour | 0.0 | 81.36 | 27.77 |
| 3 | A.Flovur | 0.0 | 68.97 | 13.97 |
| 4 | Fusurium Rigidusculum | 79.69 | 12.22 | 52.49 |
| 5 | Foxysporium. | 72.34 | 14.53 | 57.94 |
| 6 | Phoma Herbarium | 75.09 | 00 | 69.40 |
| 7 | P.Humicola | 69.28 | 00 | 55.97 |
| 8 | Drechstlera Biseptalta | 15.58 | 6.14 | 59.76 |
| 9 | Curvularia Lunata | 18.11 | 13.04 | 36.50 |
| 10 | Nigrospora Orizae | 38.29 | 0.0 | 18.30 |
| 11 | N.Sphaerica | 0.0 | 13.63 | 36.50 |
| 12 | Colletotridium Dematium | 16.25 | 0.0 | 78.34 |

Acknowledgment

We are thankful to the head, Dept of Botany P.G.College Satna & Principal(M.P)for laboratory facilities

References

1. Pandey A.K.J. Mishra, R.C Rajak, and S.K.Hasija (1996). in Herbal Medicines, Biodiversity and conservation strategies (Rajak,R.C.and Rai,M.K. eds). International Book Distributors,Dehradun: 104-138.
2. Pandey,A.K.(1999) In:microbial biotechnology. for sustainable Development and productivity. vol.1(R.C.Rajak eds)scientific Publishers,Jabalpur 85-105.
3. International seed Assoiation(1985).seed sci technol,13:299-573.
4. Neergaard,P.(197).Seed Pathology,(1and2)the Macmillan Press Ltd London.
5. Agrawal,G.P. and S.K. Hasija (1986).micro organism in the laboratory :a laboratory guide fir Mycology, Microbiology and plant Pathology.print house,lucknow (india),155.
6. Abdul Baki,A.A.,and J.D. Anderson(1972)in seed biology vol ii.(Ed., Kozlowski,T,t.) pp.283-315.,Acadmic Press New York.
7. Flannigam, B. and P.M.sellars (1977) Trans. Br. mycol. soc.,69:316-317.
8. Pandey,K.K.(1978).Acta.Mycologica14:143-149