

Influence of different temperature regimes and culture media on the growth and sporulation of *Alternaria brassicae* causing Alternaria blight of mustard

Abstract

Rapeseed-mustard (*Brassica juncea* L.) is the second most important oilseed commodity in India. *Alternaria* blight (*Alternaria brassicae*) is one of the devastating fungal pathogen, resulting in severe yield losses to the crop. The present study is conducted *in-vitro* to know the effect of different temperature regimes and various culture media on the mycelial growth and sporulation of *A. brassicae*. It is clearly evident from the results that all the tested temperature regimes and culture media showed variation in the colony diameter and other cultural characteristics. The temperature of 25°C significantly encouraged mycelial growth (85.00 mm) and exhibited excellent sporulation of *A. brassicae* compared to 15 and 35°C. In case of culture media, Potato Dextrose Agar was found to be significantly superior over the other media and registered maximum mycelial growth (82.00 mm) with excellent sporulation. However, Oat Meal Agar supported the minimum mycelial growth (45.00 mm) with fair sporulation. The variation in cultural characteristics (colony color, growth, appearance, and shape etc.) was also observed with various culture media.

Keywords: Temperature, culture media, rapeseed-mustard, *A. brassicae*, *Alternaria* blight

Introduction

Rapeseed-mustard is an economically immense and useful oilseed crop globally. In India, it is grown in diverse agro-climatic environments and is known as the third most important oilseed crop after soybean and palm [10], contributing 25 per cent of the total oilseed production [23]. The crop is a vital source of edible oil and is also used as vegetables, condiments, livestock feed, and soil amendment. Mustard oil also has demand in industries for manufacturing various products like soap, paints, hair oil, and many other products. It contains ample amounts of erucic acid, linoleic acid, and linolenic acids, which are absent in many other edible oils [21]. India occupies the second position in terms of area, covering 6.23 mha, contributing 9.43 MT annual production, and an average productivity of 1499 kg/ha [1]. The major mustard-producing states are Rajasthan (46.06%), Haryana (12.60%), and Madhya Pradesh (11.38%), which collectively contributes about 70% of the total production in India [1].

The quantity and quality of the crops is affected by various biotic and abiotic constraints. The crop suffers from various diseases, with more than 30 diseases reported in India, which not only deteriorates the seed quality but also results in a significant reduction in

oil content [18]. Amongst the fungal diseases, Alternaria blight (*Alternaria brassicae*, *A. brassicicola*, *A. raphani* and *A. alternata*) is one of the world's most catastrophic disease. Out of the four species, *Alternaria brassicae* and *A. brassicicola* are the major leaf pathogens which are known to infect the crop frequently. The severe attack of these pathogens affects not only the seed quality but also reduces the oil content significantly in different oil-yielding *Brassica* species [8, 12]. The disease causes significant loss in mustard and much prevalent in Uttar Pradesh [12]. Under favorable conditions, the losses may be as high as up to 47 per cent [17].

The disease affects all aerial parts of the plants, such as stem, leaves, fruits, and siliquae. The initial symptoms start with the appearance of black spots on leaves, stem, and siliquae which later enlarges and convert into conspicuous round spots with concentric rings exhibiting a target board-like appearance. In severe infection, many small spots conjoin to form large spots, resulting in blighting and defoliation of leaves. In a few *Brassica* species, prominent concentric rings and yellow halo are formed around the lesions [19]. The pathogen is seed as well as soil-borne in nature and can survive in soil as saprophytes on decaying plant debris.

A. brassicae is considered a pseudo-fungi with no distinguishable sexual phase and reports as cosmopolitan. Conidia are large obclavate, olive grey to dark color with longitudinal and transverse septa [2]. Crucial analysis and ample knowledge of alimentary patterns and factors influencing the growth of the pathogen (like temperature, pH, etc.) are essential for any study to understand the host-pathogen relationship. Hence, the present study is conducted *in-vitro* to know the effect of different temperature regimes and various culture media on the mycelial growth and sporulation of *A. brassicae*.

Material and methods

I. Effect of different temperature regimes on the growth of *A. brassicae*

Potato dextrose agar medium was used as a basal medium to check the disparity in radial growth and sporulation of *A. brassicae* at five different temperature regimes with an interval of 5°C, i.e., 15, 20, 25, 30, and 35°C. Inoculation was done with a 5 mm mycelial bit, cut from the seven days-old culture of *A. brassicae*, and placed in the Petri plates of 90 mm diameter consisting of sterilized solidified PDA. These Petri plates were then kept apart for incubation in various BOD incubators operating at temperatures 15, 20, 25, 30, and 35°C. These Petri plates were routinely checked after 48 h for the mycelial growth initiation of the fungus. Each treatment was replicated three times, and mycelial growth and sporulation were

recorded after seven days of inoculation in each Petri plate kept at varying temperatures [4]. The observations of the radial growth were made by drawing two perpendicular lines passing through the center of the lower surface of the bottom of the Petri plate, and the average of three replicates was assessed.

II. Effect of different culture media on the growth of *A. brassicae*

The present experiment was laid out to know the effect of eight different nutrient media, viz., Oat Meal Agar, Nutrient Agar, Richard's Synthetic Agar, Czapek's Dox Agar, Czapek's Yeast Extract Agar, Rose Bengal Agar, Corn Meal Agar, and Potato Dextrose Agar on the growth of *A. brassicae*. The media were prepared by the standardized method and autoclaved at 121.6°C, 15 lbs p.s.i. for 15 minutes. Uniform quantities (20 ml) of each culture media were poured into 90 mm diameter Petri plates. Thereafter, each Petri plate was inoculated with a 5 mm disc of a 4-5 days old culture of *A. brassicae* with the help of a sterilized cork borer. Each treatment was replicated thrice, and the Petri plates were incubated at 25±2°C for seven days. These Petri plates were frequently examined for the radial growth of the fungus [4]. The cultural characteristics (colony color, growth, margin, zonation, appearance, shape, sporulation) of all media were also recorded.

Result and Discussion

I. Effect of different temperature regimes on growth and sporulation of *A. brassicae*

The temperature ranges vary with the microorganisms as well as for host-pathogen interaction. It is evident from the results (Table 1, Plate 1) that the pathogen grew well at all the temperature regimes varying from 15 to 35°C. Significantly highest mycelial growth (85.00 mm) with excellent (++++) sporulation was observed at 25°C, which was followed by 30°C (68.66 mm), 20°C (47.33 mm), and 35°C (38.00 mm) with good (+++) to fair (++) sporulation, respectively. However, the least mycelial growth was recorded at 15°C, having poor (+) sporulation. It is revealed from the results that the most suitable temperature for the mycelial growth and sporulation of *A. brassicae* was 25°C. The results thus obtained in the current study are in agreement with the finding of other workers, who have also noted the influence

Table 1. Effect of different temperature regimes on growth and sporulation of *A. brassicae* on PDA

Temperature (°C)	Mycelial growth (mm)*	Sporulation
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15	29.66 ^e	+
20	47.33 ^c	++
25	85.00 ^a	++++
30	68.66 ^b	+++
35	38.00 ^d	++
L.S.D. (P≤0.05)	1.93	++
(A) n)±	(B)	(C)

*Each value is an average of three replicates. Values within a column followed by different alphabets are significant and some alphabets are non-significant according to Tukey's Test at P≤0.05

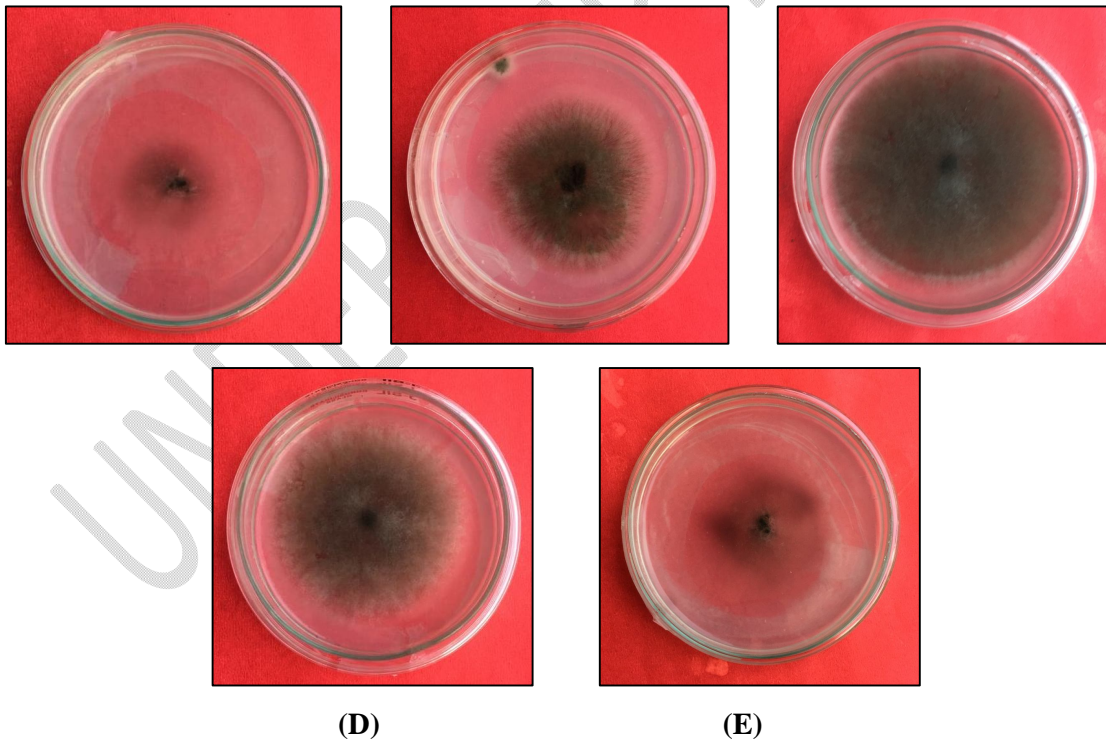


Plate 1. Effect of different temperature regimes on growth and sporulation of *A. brassicae* on PDA. 15°C (A), 20°C (B), 25°C (C), 30°C (D), and 35°C (E)

of various temperature regimes on mycelial growth and sporulation of *A. brassicae* [22, 24]. In a study, Panchal [15] recorded excellent mycelial growth and sporulation at 25°C of *A.*

(A)

(B)

(C)

alternata infecting tomato. Kurhade *et al.* [11] also observed the highest mycelial growth and excellent sporulation of *A. tagetica*, causing leaf blight of marigold at 25°C.

II. Effect of different culture media on the growth of *A. brassicae*

This experiment was carried out to note the variation in response of *A. brassicae* with respect to their cultural characteristics on eight different nutrient media. The pathogen measured significant maximum growth (82.00 mm) on Potato Dextrose Agar followed by Richard's Synthetic Agar (76.33 mm), Czapek's Yeast Extract Agar (69.66 mm), Nutrient Agar (63.66 mm), and Corn Meal Agar (61.33 mm), (Table 2). Czapek's Dox Agar was the next significantly superior nutrient medium and registered 59.00 mm growth. Minimum growth (45.00 mm) was measured on Oat Meal Agar, followed by Rose Bengal Agar (56.66 mm).

A. brassicae also showed variation in cultural characteristics on different nutrient media. The grey-color colony appeared on Oat Meal Agar and Czapek's Yeast Extract Agar, while a dark green color colony appeared on Potato Dextrose Agar. However, the rest of the tested nutrient media produced greyish white to blackish-white colony color (Table 2, Plate 2). The fluffy growth was observed in Oat Meal Agar, Richard's Synthetic Agar, Corn Meal Agar, and Potato Dextrose Agar; compressed growth was recorded in the rest of the media. The growth of the fungus was found to be fast on Potato Dextrose Agar, Richard's Synthetic Agar, Nutrient Agar, and Czapek's Yeast Extract Agar. Medium growth was found on Czapek's Dox Agar, Rose Bengal Agar, and Corn Meal Agar. While as, slow growth was observed on Oat Meal Agar (Table 2, Plate 2).

Margin, zonation, and shape also differ with the nutrient media. Round to circular margin, whereas regular to irregular shape was formed on all the tested media. Zonation in the colony was absent in all the tested media except Nutrient Agar, Czapek's Dox Agar, and Rose Bengal Agar (Table 2, Plate 2). All the tested nutrient media exhibited a wide range of sporulation. However, Potato Dextrose Agar and Richard's Synthetic Agar showed excellent (++++) sporulation. It was good (++++) on the rest of the tested nutrient media except Oat Meal Agar which exhibited fair (++) sporulation (Table 2, Plate 2).

Table 2. *In-vitro* effect of different nutrient media on mycelial growth of *A. brassicae*

S. No	Nutrient media	Mycelial growth (mm)	Colony color	Appearance	Growth	Margin	Shape	Zonation	Sporulation
1.	Oat Meal Agar	45.00 ^h	Grey	Fluffy	Slow	Rough	Irregular	Absent	++
2.	Nutrient Agar	63.66 ^d	Blackish white	Compressed	Fast	Smooth	Regular	Present	+++
3.	Richard's Synthetic Agar	76.33 ^b	Blackish White	Fluffy	Fast	Rough	Regular	Absent	++++
4.	Czapek's Dox Agar	59.00 ^f	Greyish White	Compressed	Medium	Rough	Irregular	Present	+++
5.	Czapek's Yeast Extract Agar	69.66 ^c	Grey	Compressed	Fast	Rough	Irregular	Absent	+++
6.	Rose Bengal Agar	56.66 ^g	Greyish White	Compressed	Medium	Smooth	Regular	Present	+++
7.	Corn Meal Agar	61.33 ^e	Greyish White	Fluffy	Medium	Smooth	Regular	Absent	+++
8.	Potato Dextrose Agar	82.00 ^a	Dark Green	Fluffy	Fast	Smooth	Regular	Absent	++++
L.S.D(P≤0.05)		1.86	-	-	-	-	-	-	-
S.E(m)±		0.62	-	-	-	-	-	-	-

*Each value is an average of three replicates. Values with in a column followed by different alphabets are significant and same alphabets are non-significant according to Tukey's Test at P≤0.05

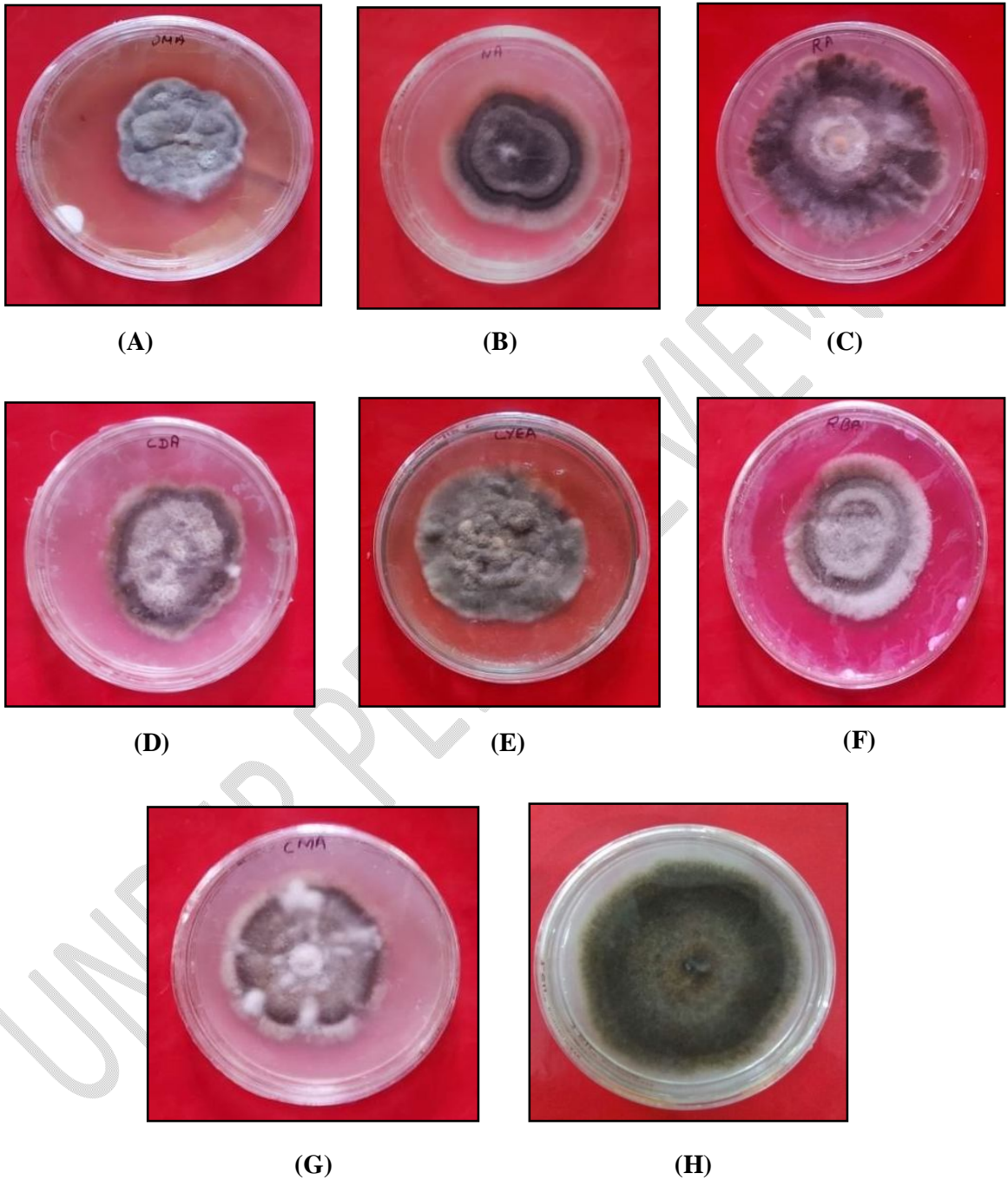


Plate 2. Growth of *A. brassicae* on different nutrient media. Oat Meal Agar (A), Nutrient Agar (B), Richard's Agar (C), Czepek's Dox Agar (D), Czepek's Yeast Extract Agar (E), Rose Bengal Agar (F), Corn Meal Agar (G), Potato Dextrose Agar (H)

The present study is in conformity with the results of Charith *et al.* [3], who studied the growth of the fungus on different media and revealed that Potato Dextrose Agar medium significantly supported the maximum radial growth (68.40 mm) of *A. brassicae* followed by Oat Meal Agar medium and the less growth was observed in the Czapek's Dox Agar and Corn Meal Agar medium. The other studies done by Nagrale *et al.* [14], Meena *et al.* [13], Koley and Mahapatra [7], Gholve *et al.* [4], Krishna *et al.* [9], and Reddy *et al.* [16] were also supported the present findings. Similar to this, Gholve *et al.* [4] observed variations in cultural features of *A. carthami* on different media and excellent sporulation on Potato Dextrose Agar. Savitha *et al.* [20] also observed variation in the colony color of isolates of *Alternaria* spp. varied from light grey to dark brown. Thus, the present investigations are in consonance with those reported earlier by several workers [5, 6, 25]. The results of the current study clearly revealed that physiological variations existed in *A. brassicae* in terms of their mycelial growth and cultural characters (colony color i.e. grey to dark green, fluffy to compressed appearance, slow to fast growth, smooth to rough margin, regular to irregular shape etc.) on various temperatures and culture media. Variations in cultural characteristics of *A. brassicae* on different culture media also provide significant knowledge that may aid in taxonomic identification.

Conclusion

The present study stated that the most appropriate temperature for growth and spore production was 25°C. On the contrary, the fungus did not grow well at 15°C, resulting in poor sporulation. In addition, among the eight tested culture media, the highest mycelial growth and excellent sporulation of pathogen were observed with Potato Dextrose Agar. However, Oat Meal Agar was found to be the least effective and supported minimum mycelial growth with fair sporulation. The study will be helpful in understanding the growth and reproduction of *A. brassicae* and its management to combat the disease caused by it. Such research might advance our understanding of the taxonomic behaviour of the fungus. In addition, the utilization of various culture media will be helpful in investigating the growth parameters of the pathogen.

Conference disclaimer:

Some part of this manuscript was previously presented in the conference: 3rd International Conference IAAHAS-2023 "Innovative Approaches in Agriculture, Horticulture & Allied Sciences" on March 29-31, 2023 in SGT University , Gurugram, India. Web Link of the proceeding: <https://wikifarmer.com/event/iaahas-2023-innovative-approaches-in-agriculture-horticulture-allied-sciences/>

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