Use of nitrogen and PGPRs for management of spot blotch in barley (*Hordeum vulgare*)

MOHAMMAD AMIR¹*, J P SRIVASTAVA², ANAM KHAN³, MANEESH KUMAR⁴, RISHABH KUMAR SINGH⁵ and F A KHAN⁶

Lovely Professional University, Phagwara, Punjab 144 402, India

Received: 28 August 2022; Accepted: 23 March 2023

Keywords: Azotobacter; Barley, Management, Spot bloch

Barley (Hordeum vulgare L.) is a widely grown crop in India and ranks 4th among cereal grains globally after wheat, rice and maize. However, its productivity in India is still below the world average. Spot blotch caused by the Bipolaris sorokiniana is one of the most important fungal diseases of barley, which causes economic injury levels with a great magnitude (Arabi et al. 2011, Al-Sadi 2016). It becomes most serious under hot and humid areas where wheat and barley are grown (Gupta et al. 2018). The seeds produced under such conditions are the main source of inoculum that exhibits poor seed germination and gives rise to diseased and frail seedlings (Neupane et al. 2010, Harding 2011). The proliferation of this disease can cut the yield as high as 30% (Singh et al. 2009, Kumar et al. 2020). Use of fungicides can have adverse effects on the environment and on consumers of crop products (Bacmaga et al. 2016). Therefore, present study was conducted to minimize the incidence of spot blotch in barley using different nitrogen levels and plant growth-promoting rhizobacteria (PGPRs).

An experiment was conducted at Research Station Masodha, Acharya Narendra Deva University of Agriculture and Technology, Kumarganj, Ayodhya, Uttar Pradesh on barley during winter (*rabi*) season of 2016–17 with three nitrogen levels (N_1 , 50%; N_2 , 75% and; N_3 , 100% and 5 PGPRs (B_1 , *Azotobacter*; B_2 , Phosphate solubilizing bacteria (PSB); B_3 , *Azotobacter* + PSB; B_4 , Biomix and; B_5 , Control) as seed treatment and replicated thrice under split plot design. Disease was allowed to develop from natural inoculums and

¹Lovely Professional University, Phagwara, Punjab; ²Acharya Narendra Deva University of Agriculture and Technology, Kumarganj, Ayodhya, Uttar Pradesh; ³Sardar Vallabhbhai Patel University of Agriculture and Technology, Meerut, Uttar Pradesh; ⁴Sardar Bhagat Singh Degree College, Goluwala, Rajasthan; ⁵Chandra Shekhar Azad University of Agriculture and Technology, Kanpur, Uttar Pradesh; ⁶Sher-e-Kashmir University of Agricultural Sciences and Technology Srinagar, Jammu and Kashmir. *Corresponding author email: drkhanfa1966@gmail.com disease rating was done by taking the per cent blighted area on flag leaf (F) and flag 1 (F-1) leaf using the rating scale as: 0, No infection; 1, up to 10%; 2, 11–20%; 3, 21–30%; 4, 31–40%; 5, 41–50%; 6, 51–60%; 7, 61–70%; 8, 71–80% and; 9, >80% leaf area blighted (Singh and Kumar 2005). The disease scores were recorded by following Kumar *et al.* (1998) using double digit scale (Table 1).

Disease severity was assessed by determining the number of lesions per cm². First and second value, respectively represents per cent blighted area on the top (flag) and second top leaf. Value 1, 2, 3, 4, 5, 6, 7, 8, and 9, respectively correspond to 10, 20, 30, 40, 50, 60, 70, 80, and 9: > 80% blighted area. The per cent disease intensity (PDI) and plant disease control (PDC) were calculated using equation I and II while Abdul Baki and Anderson (1973) were followed for seedling vigour index-I (SVI-I).

$$PDC = \frac{\text{treated plants}}{PDI \text{ in control plants}} \times 100$$
(II)

Table 1 Disease scores recorded using double digit scale based on per cent blighted area on the flag leaf and one leaf (Kumar *et al.* 1998)

Severity		Rating		
Top (flag) leaf	Second top leaf	Disease response	Range	
0	0-1	Immune (I)	00–01	
1–2	2–4	Resistant (R)	12–24	
3–4	4–6	Moderately resistant (MR)	34-46	
5-6	6–8	Moderately susceptible (MS)	56-68	
7–8	8–9	Susceptible (S)	78–89	
9	9	Highly susceptible (HS)	99	

The minimum disease severity (28.1%) with reduction of 29.5% was recorded in N₁ against the maximum disease severity (30.3%) with 24.2% reduction in N₃ (Table 2). This might be due to the negative effects of nitrogen on physical defence and the development of anti-microbial phytoalexins (Sharma 2020). Among PGPRs, the minimum disease severity (19.5%) with 51.2% reduction was recorded with B₃. This supports the findings of Biswas *et al.* (2015). The interactions of N and PGPRs showed that N₁×B₃ showed minimum disease severity (18.7%) with 53.2% reduction compared to control (N₃×B₅) and conforms the findings of Biswas *et al.* (2015). The per cent seed infection under all the N levels varied significantly (Table 2) where minimum and maximum infection (17.7 and 21.9%) was recorded with N₁ and N₃, respectively. Among PGPRs the minimum (7.2%) and maximum (34.3%) seed infection was recorded in B₃ and B₅, respectively. Reduced diseases severity due to seed treatment with *Azotobacter* has also been reported by Biswas *et al.* (2015). The interaction of N and PGPRs revealed that N₁×B₃ recorded the minimum seed infection (6.5%) compared to maximum infection (36.3%) in N₃×B₅. The result corroborates the findings of Biswas *et al.* (2015).

Table 2 Effect of nitrogen level and seed treatment with PGPRs on disease intensity and seed yield attributes of barley infected with spot botch

Treatment	PDI (%)	PDC (%)	Seed infection (%)	1000-seed weight (g)	Yield kg/ plot	Yield loss (%)
Nitrogen level	(70)	(70)	(70)	weight (g)	pior	(70)
N ₁	28.1 (31.8)	29.5 (31.6)	17.7 (24.0)	40.2	2.5	28.7 (31.2)
N ₂	28.7 (32.2)	28.1 (30.6)	19.9 (25.7)	39.5	2.4	24.8 (28.3)
N ₃	30.3 (33.2)	24.2 (25.9)	21.9 (27.1)	38.7	1.9	21.8 (24.5)
SEm±	0.08 (0.06)	0.21 (0.14)	0.20 (0.16)	0.12	0.01	0.67 (0.5)
CD (P=0.05)	0.33 (0.22)	0.84 (0.55)	0.57 (0.45)	0.33	0.04	2.70 (2.04)
Seed treatment with PGPRs						
B ₁	22.1 (28.0)	44.7 (42.0)	11.1 (19.4)	41.4	2.2	32.9 (35.9)
B ₂	35.5 (36.5)	11.2 (19.4)	28.2 (32.0)	37.6	1.8	07.9 (20.1)
B ₃	19.5 (26.2)	51.2 (45.7)	7.2 (15.5)	43.5	2.3	42.1 (43.5)
B ₄	30.0 (33.2)	24.9 (29.9)	18.4 (25.3)	39.7	2.0	20.7 (29.8)
B ₅	38.2 (38.1)	4.4 (9.8)	34.3 (35.8)	35.2	1.6	3.5 (8.7)
SEm±	0.16 (0.11)	0.41 (0.33)	0.25 (0.20)	0.15	0.01	0.57 (0.45)
CD (P=0.05)	0.48 (0.32)	1.20 (0.96)	0.74 (0.58)	0.43	0.03	1.68 (1.33)
Interaction effect						
$N_1 \times B_1$	21.7 (27.8)	45.6 (42.4)	10.0 (18.4)	42.4	2.3	43.1 (41.0
$N_1 \times B_2$	34.6 (36.0)	13.3 (21.4)	24.8 (29.8)	38.8	1.8	13.2 (21.8)
$N_1 \times B_3$	18.7 (25.6)	53.2 (46.8)	6.5 (14.8)	44.4	2.4	49.4 (44.2)
$N_1 \times B_4$	28.5 (32.3)	28.6 (32.3)	14.3 (22.2)	39.9	2.1	32.0 (34.4)
$N_1 \times B_5$	37.1 (37.5)	7.0 (15.2)	32.8 (34.9)	35.6	1.7	6.6 (14.9)
$N_2 \times B_1$	21.8 (27.8)	45.4 (42.4)	11.3 (19.6)	41.3	2.2	35.9 (36.8)
$N_2 \times B_2$	35.4 (36.5)	11.4 (19.8)	28.8 (32.4)	37.6	1.8	12.4 (20.6)
$N_3 \times B_3$	19.3 (26.0)	51.7 (46.0)	7.0 (15.3)	43.3	2.3	47.5 (43.3)
$N_4 \times B_4$	29.6 (33.0)	25.8 (30.5)	18.8 (25.6)	39.7	2.0	24.7 (29.8)
$N_5 \times B_5$	37.5 (37.7)	6.1 (14.2)	33.8 (35.5)	35.6	1.7	4.0 (11.1)
$N_3 \times B_1$	22.7 (28.4)	43.2 (41.1)	12.0 (20.3)	40.4	2.1	34.3 (35.8)
$N_3 \times B_2$	36.4 (28.4)	8.8 (17.2)	31.0 (33.8)	36.4	1.8	10.2 (18.5)
$N_3 \times B_3$	20.5 (26.9)	48.6 (44.2)	8.0 (16.4)	42.6	2.3	46.8 (42.8)
$N_3 \times B_4$	31.8 (34.3)	20.4 (26.8)	22.3 (28.1)	39.4	1.9	18.3 (25.3)
$N_3 \times B_5$	39.9 (39.2)	0.00	36.3 (37.0)	34.5	1.6	0.000
SEm±	0.13 (0.09)	0.33 (0.25)	0.45 (0.35)	0.26	0.01	0.62 (0.48)
CD (P=0.05)	0.41 (0.27)	1.04 (0.78)	1.27 (1.00)	0.74	0.04	2.25 (1.73)

Refer fo the Methodology for treatment details. Data given in parenthesis are angular transformed values.

The highest 1000-seed weight (40.2 g) and yield (2.5 kg/plot) with 28.2% increase was recorded in N₁ compared to a lower 1000-seed weight (39.5 g) and yield (2.4 kg/plot) with 24.8% increase in N₂. A higher 1000-seed weight under lower dose of N may be attributed to declined disease infection under lower N dose in the present study. Among PGPRs the maximum 1000-seed weight (43.5 g) and yield (2.3 kg/plot) with 42.1% increase was recorded in B₃ followed by B₁ (41.4 g 1000-seed weight and 2.28 kg/plot yield with 32.9% yield increase) against the minimum 1000-seed weight (35.2 g) and yield (1.6 kg/plot) with control (B₅) (Table 2). Among different interactions, the maximum 1000-seed weight (44.4 g) and yield (2.4 kg/plot) with 49.4% increase was recorded with N₁×B₃ over control.

Table 3 clarified that the highest seed viability (70.0%), germination (67.0%), seedling length (23.7 cm) and SVI-I (1663.0) was recorded in N₁ compared to minimum seed vigour (62.0%), germination (57.0%), seedling length (21.8 cm) and SVI-I (1329.8) in N₃. Regarding PGPRs, the maximum seed viability (93.0%), germination (90.0%), seedling length (27.2 cm) and SVI-I (2459.7) was evident in B₃. However, the interaction of N and PGPRs showed significant effect only with seedling length and the maximum seedling length (28.4 cm) being in N₁×B₃ followed by 27.2 cm in N₂×B₃ over the control (16.2 cm). The present findings are in accordance with the findings of Singh and Kumar (2008).

Table 3	Effect of nitrogen level and	l seed treatment with PGPRS on a	seed quality of barley	infected with spot botch

Treatment	Viability (%)	Germination (%)	Seedling length (cm)	SVI-I
Nitrogen level				
N ₁	70.0 (59.0)	67.0 (56.0)	23.7	1663.0
N ₂	67.0 (56.0)	62.0 (53.0)	22.9	1491.2
N ₃	62.0 (53.0)	57.0 (50.0)	21.8	1329.8
SEm±	0.53 (0.36)	0.67 (0.45)	0.14	18.32
CD (P=0.05)	1.046	1.30	0.41	53.15
Seed treatment with PGPRs				
B ₁	82.0 (65.0)	79.0 (63.0)	25.1	1978.8
B ₂	51.0 (45.0)	46.0 (43.0)	20.6	962.1
B ₃	93.0 (75.0)	90.0 (02.0)	27.2	2459.7
B ₄	68.0 (56.0)	64.0 (53.0)	23.8	1531.4
B ₅	37.0 (37.0)	31.0 (34.0)	17.2	541.4
SEm±	0.69 (0.47)	0.87 (0.58)	0.18	23.6
CD (P=0.05)	1.35	1.67	0.52	68.6
Interaction effect				
$N_1 \times B_1$	85.0 (68.0)	84.0 (66.0)	25.5	2,119.2
$N_1 \times B_2$	55.0 (48.0)	52.0 (46.0)	22.1	1168.5
$N_1 \times B_3$	96.0 (78)	92.0 (73.0)	28.4	2616.3
$N_1 \times B_4$	74.0 (59.0)	71.0 (58.0)	24.4	1735.5
$N_1 \times B_5$	42.0 (40.0)	38.0 (38.0)	17.8	675.6
$N_2 \times B_1$	82.0 (65.0)	79.0 (63.0)	25.3	2001.6
$N_2 \times B_2$	53.0 (47.0)	45.0 (42.0)	21.0	944.4
$N_3 \times B_3$	93.0 (75.0)	91.0 (72.0)	27.2	2472.4
$\mathbf{N_4}\times\mathbf{B_4}$	68.0 (56.0)	64.0 (53.0)	23.4	1495.9
$N_5 \times B_5$	37.0 (38.0)	31.0 (34.0)	17.4	541.6
$N_3 \times B_1$	78.0 (62.0)	74.0 (60.0)	24.5	1815.6
$N_3 \times B_2$	45.0 (42.0)	41.0 (40.0)	18.8	773.4
$N_3 \times B_3$	91.0 (72.0)	88.0 (70.0)	26.0	2290.3
$N_3 \times B_4$	63.0 (53.0)	58.0 (50.0)	23.5	1362.8
$N_3 \times B_5$	32.0 (34.0)	25.0 (30.0)	16.2	407.1
SEm±	01.19 (0.81)	1.45 (0.66)	0.31	40.95
CD (P=0.05)	(NS)	NS	0.91	NS

Refer to the Methodology for treatment details. NS, Non-significant. Data given in parenthesis are in angular transformation.

SUMMARY

Spot blotch disease of barley caused by Bipolaris sorokiniana is prevalent everywhere but causes significant yield losses under warm and humid climates. Use of fungicides is a common practice to control the spot blotch but at the same time it can pose a risk to environment as well as humans consuming such products. A field experiment was conducted to minimize the incidence of spot blotch in barley using different nitrogen levels and PGPRs as seed treatment. Application of lower nitrogen dose (50% of RDF $-N_1$) and seed treatment with Azotobacter + PSB $-B_3$) either individually or in combination significantly reduced the severity of spot blotch and increased the yield of barley. As such it may be concluded that spot blotch disease of barley can be managed by use of lower dose of nitrogen along with seed treatment with Azotobacter + PSB and use of fungicide may be avoided.

REFERENCES

- Abdul Baki A A and Anderson J D. 1973. Vigour determination in soybean by multiple criteria. *Crop Science* **13**(6): 630–33.
- Al-Sadi A M. 2016. Variation in resistance to spot blotch and the aggressiveness of *Bipolaris sorokiniana* on barley and wheat cultivars. *Journal of Plant Pathology* 98(1): 97–103.
- Arabi M I E, Al-Daoude A, Shoaib A and Jawhar M. 2011. Transcriptional interactions during barley susceptible genotype infection with *Cochliobolus sativus*. *Russian Journal of Genetics* 47(7): 879–83.
- Bacmaga M, Wyszkowska J and Kucharski J. 2016. The effect of the Falcon 460 EC fungicide on soil microbial communities, enzyme activities and plant growth. *Ecotoxicology* 25(8): 1575–87
- Biswas S K, Shankar U, Kumar S, Kumar A, Kumar V and Lal K. 2015. Impact of bio-fertilizers for the management of spot blotch

disease and growth and yield contributing parameters of wheat. *Journal of Pure and Applied Microbiology* **1**(4): 3025–31.

- Gupta P K, Chand R, Vasistha N K, Pandey S P, Kumar U, Mishra V K and Joshi A K. 2018. Spot blotch disease of wheat: The current status of research on genetics and breeding. *Plant Pathology* 67(3): 508–31.
- Harding H. 2011. Effect of *Bipolaris sorokiniana* on germination and seedling survival of varieties or lines of 14 *Triticum* species. *Canadian Journal of Botany* 49(2): 281–87.
- Kumar J, Singh G and Nagarajan S A. 1998. Field scale for leaf blight recording. *Indian Wheat Newsletter* 5(2): 3–4.
- Kumar S, Kumar N, Prajapati S and Maurya S. 2020. Review on spot blotch of wheat: An emerging threat to wheat basket in changing climate. *Journal of Pharmacognosy and Phytochemistry* 9(2): 1985–97.
- Neupane A C, Sharma R C, Duveiller E and Shrestha S M. 2010. Sources of *Cochliobolus sativus* inoculum causing spot blotch under warm wheat growing conditions in South Asia. *Cereal Research Communication* 38: 541–49.
- Sharma S. 2020. Impacts of nitrogen on plant disease severity and plant defence mechanism. *Fundamental and Applied Agriculture* **5**(3): 303–14.
- Singh D P and Kumar P. 2005. Method of scoring leaf blight of wheat caused by *Bipolaris sorokiniana* (Sacc.) Shoem. on top leaves at adult plant stage. *Integrated Plant Disease Management*, pp. 289–54. Sharma R C and Sharma J N (Eds.), Scientific Publishers, Jodhpur, India.
- Singh D P and Kumar P. 2008. Role of spot blotch (*Bipolaris sorokiniana*) in deteriorating seed quality, its management in different wheat genotypes using fungicidal seed treatment. *Indian Phytopathology* **61**(1): 49–54.
- Singh R N, Singh A K and Singh S P. 2009. Prevalence and management of spot blotch (*Cochliobolus sativus*) of barley (*Hordeum vulgare* L.) in eastern India. Proceedings of the National Academy of Sciences India. Section B - Biological Sciences **79**(1): 65–69.