

Journal of Advances in Biology & Biotechnology

Volume 27, Issue 11, Page 603-613, 2024; Article no.JABB.125806 ISSN: 2394-1081

Innate Immunity in Rice: A Defense Against Bacterial Leaf Blight

Sushitha ^{a*} and A. Sajeena ^b

 ^a Kerala Agricultural University, Department of Plant Pathology, College of Agriculture, Vellayani, Thiruvananthapuram 695 522, India.
 ^b Kerala Agricultural University, Integrated Farming System Research Station, Thiruvananthapuram 695 002, India.

Authors' contributions

This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.

Article Information

DOI: https://doi.org/10.9734/jabb/2024/v27i111644

Open Peer Review History: This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: https://www.sdiarticle5.com/review-history/125806

Review Article

Received: 03/09/2024 Accepted: 05/11/2024 Published: 12/11/2024

ABSTRACT

Rice (*Oryza sativa* L.) is one of the major staple food crops of the world and sustainable rice production is important for ensuring global food security. Throughout the growing season, a variety of pathogens including fungi, bacteria, viruses, and nematodes, infect different parts of the crop resulting in yield loss. Bacterial leaf blight caused by *Xanthomonas oryzae* pv. *oryzae* (*Xoo*) is one of the major limiting factors in rice production. Being a vascular pathogen, *X. oryzae* pv. *oryzae* interferes with a range of physiological and biochemical exchange process in rice. The earlier the pathogen infection, more will be the loss and grain quality is also severely affected due to infection. The spread of the pathogen is too rapid in the favourable climatic conditions. The disease results in 20 to 30 per cent annual loss in rice production and under severe conditions, the yield loss goes up to 80 per cent. Plant pathogens employ different ways to attack host plants and impair plant growth and reproduction. Unlike vertebrates, plants lack mobile immune cells and an adaptive immune system. Plants mainly rely on two interconnected tiers of the innate immune system to perceive and

Cite as: Sushitha, and A. Sajeena. 2024. "Innate Immunity in Rice: A Defense Against Bacterial Leaf Blight". Journal of Advances in Biology & Biotechnology 27 (11):603-13. https://doi.org/10.9734/jabb/2024/v27i111644.

^{*}Corresponding author: E-mail: sushitha87@gmail.com;

respond to pathogen infection. This innate immune system or basal resistance mediated by a repertoire of Resistance or R genes in plants acts as the first line of pre-formed and inducible defenses that protect the host plants from large number of pathogens. Over the years, extensive investigation on the molecular interactions between rice and *Xanthomonas oryzae* pv. *oryzae* has made impressive progress in understanding the molecular basis of rice innate immunity against bacterial leaf blight. Improving plant immunity has been considered as one of the best choices available for achieving economical and sustainable management of bacterial leaf blight in a durable manner. In this review, we summarize the molecular basis of two tiered innate immune system including PAMP triggered immunity (PTI) and Effector triggered immunity (ETI) as well as R genes involved in rice - *Xanthomonas oryzae* pv. *oryzae* p

Keywords: Xanthomonas oryzae pv. oryzae; bacterial leaf blight; r genes; innate immune system; defense.

1. INTRODUCTION

"Rice is the most important cereal crop in the world feeding more than 50 per cent of the world population. It is a rich source of carbohydrates and energy and provides 20 per cent of the world calories requirement" (Khush 2005). "Demand for rice is increasing year after year with the increase in population. But rice production is limited by various biotic and abiotic factors. Among them, vulnerability of the crop to diseases results in drastic crop loss and yield reduction. A large number of fungi, bacteria and virus infect rice crop. Bacterial leaf blight (BLB) of rice is one of the most devastating and economically important diseases in rice. It is caused by the bacterial pathogen **Xanthomonas** orvzae pv. orvzae, which has been ranked as the fourth most important plant pathogenic bacteria worldwide" (Mansfield et al. 2012). "The disease causes yield loss in the range of 20 to 30 per cent, but under severe infection, yield loss up to observed" 80 has per cent been (Gnanamanickam et al. 1999). "Several control measures such as chemical and biological methods are used to control the spread of the disease. Unfortunately, these measures could not effectively control the disease. The use of pesticides is expensive and are not eco-friendly and biological control alone cannot completely control the disease. Therefore, one of the best, economic and safe ways to manage this disease is by boosting the innate resistance of the crop against pathogen attack" (Pradhan et al. 2015).

"Microbes live in nearly every environment and even in nutrient limited conditions in soil and water. Eukaryotes provide an ideal habitat for various microbes. Even though plants and animals are surrounded by millions of microorganisms, they remain healthy in most of the situations. Plants are attacked by large number of microbes, but only few succeed in causing disease. Others are detected by multiple layers of sophisticated surveillance mechanisms that recognize potentially dangerous pathogens and rapidly respond before those organisms have a chance to cause serious damage. This is known as the innate immunity or basal resistance of the plants and it is mediated by a repertoire of Resistance or R genes in plants. By using this innate immunity, plants encounter majority of the pathogens which come in their way. Hence, one of the best ways to manage BLB is by enhancing the innate immune system of rice by breeding for resistance varieties" (Mundt 2014).

2. BACTERIAL LEAF BLIGHT OF RICE

"Bacterial blight is a seed borne disease caused by the gram-negative bacterium, X. oryzae pv. orvzae. It is a severe threat to rice production. The disease was first observed by the farmers of the Fukuoka area of Japan, in 1884" (Mizukami and Wakimoto 1969). "In India, bacterial leaf blight disease incidence was first reported in Maharashtra" (Srinivasan et al. 1959). "Earlier, the disease was considered to be of minor importance in India until it broke out in an epidemic form in Shahabad district of Bihar in 1963. This disease can affect rice plants at any plant growth stages. Bacterial leaf blight generally causes yield loss ranging from 20 to 30 per cent. In case of severe infection, disease cause yield loss up to 80 per cent besides severely affecting the grain quality" (Gnanamanickam et al. 1999).

3. PATHOGEN MODE OF INFECTION AND SYMTOMATOLOGY

"X. oryzae pv. oryzae enters the rice leaf typically through hydathodes at the leaf tip and leaf margin" (Ou 1985). "Subsequently, it multiplies and flourishes in the intercellular spaces (apoplast) beneath the epithelial cells and later disseminates to other parts of plants through the xylem vessels" (Noda and Kaku 1999). "Within a few days of infection, bacterial cells and Exopolysaccharides (EPS) fill the xylem vessels and ooze out from hydathodes, forming beads or strands of exudate on the leaf surface, a characteristic sign of the disease and a source of secondary inoculum" (Mew et al. 1993).

"The symptom of bacterial leaf blight is first observed as water-soaked spots at the leaf tips and margins. The spots expand along the veins, merge, and become chlorotic and then necrotic, forming opaque, white to grey coloured lesions that extend from the leaf tip down along the leaf veins and margins. Symptoms are often evident at the tillering stage and become severest at the flowering stage. In the tropics, susceptible rice plants show a seedling blight disease called Kresek. which is а more destructive manifestation of the disease in which the entire foliage becomes pale yellow and wilts, resulting in a partial or total failure of the crop" (Niño-Liu et al. 2006).

4. INNATE IMMUNE SYSTEM OF PLANTS

"Immune system is essential for the survival of all living organisms. Without an immune system, both plants and animals would be open to attack to many microbes such as bacteria, viruses, fungi and protozoa. Plants and animals should avoid becoming a meal to microbes, which vastly outnumber eukaryotic life in both quantity and diversity. It is the immune system that keeps the plants and animals safe and healthy against a sea of pathogens. Both plants and animals have an innate immune system, which protect them from the majority of microbes they encounter during their lifetime" (de Wit 2007, Haney 2014).

"Plants have multiple layers of sophisticated surveillance mechanisms that recognize dangerous pathogens and rapidly respond before those organisms which have a chance to cause damage. serious These surveillance mechanisms are linked to some specific preprogrammed defense responses and this is known as the innate immunity or the basal resistance of the plant. It is the first line of preformed and inducible defense that protect against entire groups of pathogens. In short, the innate ability to defend against pathogen attack

is known as innate immunity of the host" (Liu 2014).

Plants have developed a two-layered innate immune system that includes pathogenassociated molecular pattern (PAMP) triggered immunity (PTI) and effector triggered immunity (ETI). PTI is the first line of defense, which is governed by pattern recognition receptors (PRRs) that recognize highly conserved PAMPs to trigger a relatively weak immune response that restricts colonization by invading organisms (Ausubel 2005). In contrast, ETI, the second line of defence, is a rapid and robust response, usually associated with a hypersensitive reaction (HR) in plants. ETI is initiated by R proteins that directly or indirectly recognize highly variable pathogen produced molecules called avirulence (Avr) effectors (Kushalappa et al. 2016). The innate immune system of plants operates through three stages Viz., PAMP triggered immunity (PTI), Effector triggered susceptibility (ETS) and Effector triggered immunity (ETI) (Fig. 1).

5. PAMP-TRIGGERED IMMUNITY (PTI)

"PTI involves the recognition of conserved, indispensable microbial elicitors called microbe/pathogen associated molecular patterns by a class of plasma membrane bound extracellular receptors called PRRs" (Jones and Dangl 2006). "Activation of PRRs leads to intracellular signaling, transcriptional reprogramming, and biosynthesis of a complex output response that limits colonization, which ultimately contribute to halt infection before the pathogen gains a hold in the plant" (Muthamilarasan and Prasad 2013).

"PAMPs essential structures or components that are conserved throughout the whole classes of pathogens. This includes oligogalacturonides, ergosterol, bacterial flagellin, pep-13, xylanase, cold-shock proteins, lipopolysaccharides (LPS) etc. Many plant pathogens produce lytic enzymes to break the structural barriers of plant tissues. The products formed as a result of these enzymatic degradation such as cell wall fragments and peptides, can function as endogenous elicitors called damage-associated molecular patterns (DAMPs). These DAMPs emerge in the apoplast and serve as danger signals to induce innate immunity similar to MAMPs" (da Cunha et al. 2006).

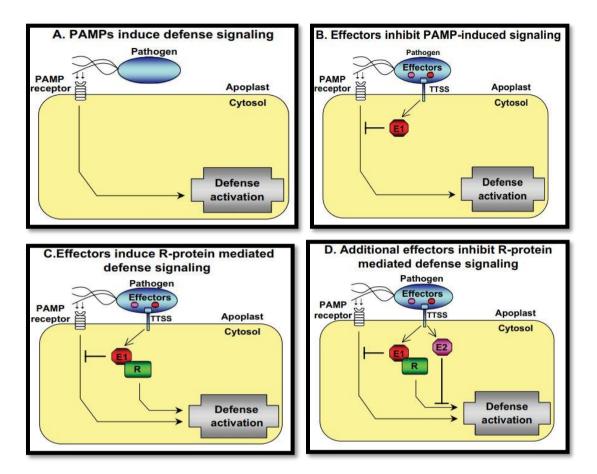


Fig. 1. Innate immune response in plants (a) PTI (b) ETS (c) ETI (d) ETS (da Cunha et al., 2006)

"PRRs are plasma membrane localised proteins capable of recognizing PAMPs and activating PTI. PRR familv The includes receptor-like kinases (RLK) and receptorand Rathjen like proteins (RLP)" (Dodds 2010).

Effector Triggered Susceptibility (ETS): "Successful pathogens have many evolved strategies to challenge the PTI and promote pathogenesis by injecting a set of effector proteins across the plant cell wall into the cytoplasm through the type III secretion system (TTSS) and this is known as effector triggered susceptibility" (Iriti and Faoro 2007).

Effector Triggered Immunity (ETI): "Effector triggered immunity (ETI) is activated by the effectors produced the pathogens. by Receptors on the plant recognize the pathogen effectors and activate ETI. FTI immune response is depended on R genes, and is activated by specific pathogen strains. Plants often cause an FTI apoptotic hypersensitive response" (Nürnberger et al. 2004).

6. RESISTANCE GENES (R GENES)

"Plants have evolved R genes whose products allow recognition of specific pathogen effectors, either through direct binding or by recognizing effector's alteration of host proteins" (Jones and Dangl 2013). "Plants encode many types of extracellular and intracellular R proteins. Many R genes encode NB-LRR type proteins (nucleotidebinding/leucine-rich repeat domains, also known as NLR proteins or STAND proteins). R gene products control a broad set of disease resistance responses whose induction is often sufficient to stop further spread or growth of the pathogen. Most of the plant immune systems carry a repertoire of 100-600 different R genes that mediate resistance to various virus, bacteria, fungus, oomycete, nematodes and insects. R genes are generally very specific to particular pathogen strains" (de Wit 2007).

7. INNATE IMMUNITY IN RICE AGAINST X. ORYZAE PV. ORYZAE

"Like other plants, rice also has evolved a twolayered innate immune system which includes PTI and ETI. R genes form the basis of this innate immune system of rice plant. To date 45 genes conferring BLB resistance have been reported in rice plant and these R genes are designated in a series from Xa1 to Xa45" (Kesh and Kaushik 2020). "Out of these, 17 genes viz., xa5 xa8, xa13, xa15, xa19, xa20, xa24, xa25, xa26, xa28, xa31, xa32, xa33, xa34, xa41(t), xa42 and xa44(t)74 are recessive and remaining genes are dominant. Out of these 45 resistance genes in rice, Xa1, Xa3/26, Xa5, Xa10, Xa13, Xa21, Xa23, Xa25, XA4, Xa41 and Xa27 have been cloned successfully and Xa2, Xa4, Xa7, Xa22, Xa30, xa31 xa33, xa34, Xa38, Xa39, Xa40, xa42 have been fine mapped. All these resistance genes follow Mendelian pattern of gene inheritance and express resistance to a diverse strains of X.oryzae pv.oryzae" (Lee et al. 2011). hese R genes identified in rice can be categorized into four groups based on the encoded proteins such as receptor kinases, SWEET genes, Executer genes and other genes (Jiang et al. 2020).

8. RECEPTOR KINASES

"PRRs, which can recognize diverse groups of PAMPs form the key component of the innate immune system of the plant. All the known plant PRRs falls into mainly two groups such as transmembrane receptor like kinases (RLKs) or transmembrane receptor like proteins (RLPs). There are over 1100 identified RLKs/RLPs in rice genome. RLKs contain an extracellular domain, a transmembrane domain, and an intracellular kinase domain, whereas RLPs lack the kinase domain. Leucine rich repeat receptor like kinases (LRR-RLKs) are the largest subfamily of plant RLKs" (Song et al. 1995).

"The rice genome encodes more than 290 LRR-RLKs. Xa21 and Xa3/Xa26 are the wellcharacterized members as well as representatives of non-RD (non-arginineaspartate) receptor kinases and both confer broad-spectrum resistance to X. oryzae pv. oryzae strains. Xa4 gene is a wall associated kinase belonging to receptor kinase family" (White and Yang 2009).

"Xa21 was the first cloned R gene in rice and it was originated from the wild rice species *Oryza longistaminata*" (Khush et al. 1990). "These proteins have an extracellular LRR domain, a transmembrane motif, a juxta membrane domain and a cytoplasmic kinase domain" (Song et al. 1995). "XA21-mediated immunity is activated

upon recognition of a 194-amino acid protein, Ax21 (activator of XA21-mediated innate immunity)" (Lee et al. 2009). "*Xa21* confer broadspectrum resistance to *X* .oryzae pv. oryzae. *Xa21* recognizes, a sulfated 17-amino acid peptide derived from the amino terminus of Ax21, a protein secreted through type 1 secretion system of *X*. oryzae pv. oryzae and thus activates PTI" (Chen et al. 2014).

9. XA21 MEDIATED DEFENSE AGAINST X. ORYZAE PV. ORYZAE

"Ax21 protein is a sulfated peptide secreted by *X. oryzae* pv. *oryzae* through the type I secretion system. This protein triggers XA21-mediated resistance by binding to the LRR domain of XA21. Because this peptide is conserved in many *Xanthomonas* species and even outside the *Xanthomonas* genus, it is considered as a PAMP molecule and XA21 as a PRR protein" (Lee et al. 2009).

"XA21 mediates a high level of PTI but with race specificity to X. oryzae pv. Oryzae" (Ji et al. 2018). "Several XA21 binding (XB) proteins have been reported to be involved in the rice defense response against X. oryzae pv. oryzae. XB24/ATPase promotes autophosphorylation of XA21 at ser/thr kinase (Ser 686, thr688, and Ser689) domains in the JM which maintains an inactive status for XA21 protein. Upon recognition of Ax21, XB24 dissociates from XA21. leading to its activation. XB3/E3 ubiquitin ligase is a substrate of XA21 kinase activity and is required for XA21-mediated resistance against BLB. XA21 transphosphorylates XB3, and activate MAPK cascade. Once the signal has been replayed, XA21 binds with the XB15 phosphates, which attenuates the immune response by dephosphorylation of aminoacids required for XA21 function. XB15/protein phosphatase 2C can dephosphorylate XA21, which results in the inactivation of XA21. The transcriptional regulator XB10/WRKY62 negatively regulates Xa21-mediated disease resistance. Cleavage XA21 of and translocalization of its kinase domain to the nucleus, where it interacts with WRKY62, is required for the Xa21-initiated defense response" (Jiang et al. 2012).

XA3/XA26 mediated defense: "Another LRR-RLK gene is *Xa26* and it was originally identified from indica variety Minghui 63, an elite restorer line of hybrid rice in China. Further studies demonstrated that, *Xa3* gene identified in a japonica variety Wase Aaikoku 3, is the same gene as that of Xa26 gene. Xa3/Xa26 mediates a race-specific resistance to many strains of X. oryzae pv. oryzae but with a resistance spectrum different from Xa21 gene mediated defense. Xa3/Xa26 locus confers a durable resistance to strains of X. oryzae pv. oryzae. A gene-dosage effect exists in Xa3/Xa26-mediated defense. Higher expression level of Xa3/Xa26 confers a stronger, wider spectrum, and whole-growthstage resistance to pathogen" (Liu et al. 2018).

XA4 mediated defense: "Xa4 gene encodes a cell wall-associated kinase and confers a racespecific resistance to X. oryzae pv. oryzae at all stages of rice growth. Wall associated kinases (WAKs) are also a subfamily of RLKs that physically link the cell wall with the plasma membrane and transmit extracellular signals to the cytoplasm. In the early 1970s, Xa4 was first introgressed into commercial rice varieties. It is one of the most widely employed resistance genes in breeding programmes in rice. Nearly all the indica hybrid rice cultivars in China carry the Xa4 gene. Xa4 prevents the invasion of X. oryzae pv. oryzae through reinforcing the plant cell wall. The Xa4 mediated resistance is associated with the accumulation of two phytoalexins, viz., sakuranetin and momilactone A, which are likely to suppress X. oryzae pv. oryzae in plants. In addition to conferring durable resistance to X. oryzae pv. oryzae, Xa4 also increases the mechanical strength of the culm and reduces the plant height slightly, and thus may enhance the lodging resistance. The multiple favourable agronomic traits related with Xa4 explain why it is widely used in breeding programmes to develop resistant varieties" (Yoshimura et al. 1998).

"Xa4 mediated defense operates through cell wall reinforcement, secretion of phytoalexins and by enhanced synthesis of jasmonic acid is (JA-Ile). Cellulose isoleucine mainly synthesized by cellulose synthase (CESA) family enzymes. Rice CesA4, CesA7 and CesA9 genes are responsible for the secondary cell wall synthesis. The expression of CesA7 and CeSA9 genes was enhanced during Xa4.mediated defense. Accumulation of phytoalexins such as sakuranetin and momilactone A, along with the increase in JA-Ile abundance occurs during Xa4 mediated defense. Xa4-mediated resistance may require production of the bioactive JA-Ile, which in turn stimulates production of phytoalexins against X. oryzae pv. Oryzae" (Hu et al. 2017).

10. SWEET GENES

"SWEET genes encode sugar transporter proteins, which are involved in exporting sugar across the membrane into the apoplast. During infection, these SWEET genes will be hijacked by transcription activator like effectors (TALE) of the pathogen, which is essential for growth and virulence of the pathogen. These SWEET proteins transport sugar across the membrane to the apoplast area which is essential for the survival of the pathogen. But any mutation in these genes confers resistance to pathogen attack. Three recessive R genes such as *xa13*, *xa25* and *xa41* represent well studied examples of SWEET genes" (Zhang and Wang 2013).

11. XA13 MEDIATED DEFENSE

"Rice plants carrying recessive xa13 exhibit specific resistance to Xoo strain PXO99" (Verdier et al. 2012). "This resistant gene is widely used for breeding programmes in South Asian countries" (Jiang 2012). "The dominant allele of the of this recessive gene, Xa13 (also named Os8N3 and OsSWEET11), is a susceptibility gene specific to PXO99 strain, which secretes the TAL effector PthXo1 during infection" (Yang et al. 2006). "When rice plants are infected with X. oryzae pv. oryzae, TAL effectors are secreted into the plant cell. The TAL effector pthXo1 injected into the nucleus directly target the effector binding element (EBEs) (UPT_{PthXol} box) in the promoter of dominant Xa13 gene and activates its expression. Xa13 encode a plasma membrane located sugar transporter gene, which is involved in the release of sugar into the apoplast and xylem, where the X. oryzae pv. oryzae resides, thus providing nutrition for the Xoo. Xa13 can also interact with two plasma membrane located copper transporter-type proteins, COPT1 and COPT5, to promote removal of copper from the xylem vessels. Copper is an essential micronutrient for plants but it inhibits Xoo growth, and PXO99 is more sensitive to copper than other Xoo strains When TAL effector activates SWEET genes, these copper transporter proteins also get activated. Copper transporters facilitate the removal of Cu from the xylem vessels and apoplast thereby increasing the virulence of the bacteria" (Yuan et al. 2010). "Resistance of rice plants carrying recessive xa13 gene is due to the mutation of UPT_{PthXol} box in xa13 promoter" (Chu et al. 2006). "The effectors secreted cannot bind to the EBE which results in PXO99 strain being unable to induce recessive xa13 expression. The copper

level in the xylem vessels of rice plants carrying recessive *xa13* can inhibit PXO99 growth and plants have passive resistance to *Xoo*. But when the gene is recessive effectors cannot bind to EBE of target genes, there by imparting resistance" (Zhang and Wang 2013).

Executer Genes: Xa27, Xa10 and Xa23 are the three executor genes in rice with multiple potential transmembrane domains functioning as a promoter trap, which are transcriptionally activated by TAL effectors and trigger defense responses. Dominant Xa27 mediated race specific resistance depends on its transcriptional activation by AvrXa27, a TAL effector of Xoo. Pathogen secreting TAL effector AvrXa27, would be directed in to the nucleus, where it interacts with the up regulated bv transcription factor (UPT) box in the promoter region of the target Xa27 gene and activates its expression leading to the resistance (Gu et al. 2005).

Other Genes: "In rice genome, 480 nucleotidebinding domain and leucine-rich repeat (NLR) genes have been revealed, but only a single one, *Xa1*, conferring resistance to *X. oryzae* pv. *oryzae*, was isolated. *Xa1* was isolated from the japonica cultivar Kogyoku and its expression was induced by bacterial infection and wounding. This gene confers race-specific resistance to *Xoo* strain T7174" (Yoshimura et al. 1998).

"The recessive gene xa5 confers broad resistance spectrum to *X. oryzae* pv. *oryzae*. The *xa5* is a recessive allele of the gene Xa5, which encodes transcription factor IIA gamma subunit 5 (TFIIA γ 5). TFIIA is a basal transcription factor of eukaryotes which is essential for polymerase II-dependent transcription. TFIIA γ 5 is hijacked by TAL effectors by direct physical interaction with a transcription factor binding (TFB) region of TALE during infection. But the recessive gene xa5 contains a mutation in the 39th residue, in which the valine (V) residue is replaced with glutamine (E). The TAL effector cannot interact with recessive genes, leading to resistance" (Jiang et al. 2012).

12. MANAGEMENT OF BACTERIAL LEAF BLIGHT

"Use of antibiotics such as streptocyclin for controlling BLB is not eco-friendly. Host plant resistance is generally the most favourable tactic to control diseases due to economic and environmental reasons. Marker-assisted selection (MAS) and genetic transformation are two major approaches for R gene the in plant breeding deployment programs. Pyramiding R genes resistant to different races through the pathogen markerof assisted breeding strategies, is a very effective broaddurable way to achieve and spectrum resistance, while employment of a R and adaption single gene of the pathogen often lead to resistance breakdown in a short period. Based on reports, xa5, Xa7, xa13, Xa21 and Xa23 are more frequently used by rice breeders due to the comparatively broader spectra of resistance" (Kumar et al. 2020).

Combination of gene	Variety/genotype	Country	Reference
xa5, xa13 and XA21	PR106	India	Singh <i>et al.</i> (2001) [44]
xa5, xa13 and XA21	Samba Mahsuri	India	Sundaram <i>et al.</i> (2008) [45]
XA21, xa13 and sd-1	Type 3 Basmati	India	Rajpurohit <i>et al</i> . (2011) [46]
<i>Xa4, xa5, xa13</i> and <i>XA21</i>	Mahsuri	India	Guvvala et al. (2013) [47]
Xa38, xa13 and XA21	PAU 201	India	Sundaram <i>et al.</i> (2014) [48]
XA21, xa13, xa5 and Xa4	Lalat and Tapaswini	India	Sundaram <i>et al.</i> (2014) [48]
XA21, xa13 and xa5	Swarna and IR64	India	Sundaram <i>et al.</i> (2014) [48]
XA21 and xa13	Pusa Basmati 1	India	Sundaram <i>et al.</i> (2014) [48]
XA21	Zhongyou 1176	China	Cao <i>et al.</i> (2003) [49]
XA21	Zhongyou 6	China	Cao <i>et al.</i> (2003) [49]
Xa4 and Xa5	Angke	Indonesia	Sattari et al. (2014) [50]
<i>Xa5, Xa13</i> and <i>Xa21</i>	Improved Pusa	India	Gopalakrishnan et al. (2008)
	Basmati-1		[51]
Xa21, xa13, and xa5	Ranidhan	India	Mohapatra et al. (2023)[52]

Table 1. Cultivars improved for bacterial blight resistance through marker assisted breeding

Tainung 82 (TNG82) is one of the most popular japonica varieties in Taiwan due to its relatively high yield and grain quality, however, TNG82 is susceptible to bacterial blight (BB) disease. The most economical and eco-friendly way to control BB disease in japonica is through the utilization of varieties that are resistant to the disease. In order to improve TNG82's resistance to BB disease, five bacterial blight resistance genes (Xa4, xa5, Xa7, xa13 and Xa21) were derived from a donor parent, IRBB66 and transferred into TNG82 via marker-assisted backcrossing breeding. These individuals displayed a high level of resistance against the bacterial leaf blight. The five identified bacterial leaf blight gene pyramided lines exhibited yield levels and other desirable agronomic traits, including grain quality and palatability, consistent with TNG82. Bacterial blight resistant lines possessing the five identified leaf blight genes exhibited not only higher levels of resistance to the disease, but also greater yield levels and grain quality (Yu-Chia et al., 2020).

13. CONCLUSION

Bacterial leaf blight disease caused by X. oryzae pv. oryzae is one of the most widespread and economically important diseases of rice. It is a threat to rice production in tropical and temperate countries due to its high epidemic potential. X. oryzae pv. oryzae has been ranked as one among the top ten economically important bacterial plant pathogens worldwide. Bacterial leaf blight is very difficult to manage and none of the methods could completely manage the problem. One of the best ways to manage the disease is by boosting the innate resistance of the host plant against the pathogen. It can be achieved by incorporating resistant genes in cultivars. It has been proven that adoption of resistance is the most effective, economic, and environment-friendly strategy to avoid vield loss caused by bacterial leaf blight disease. During the co evolution of rice with pathogen, successful defense systems with the core of R genes have been evolved by rice to resist the disease. These R genes can be utilized in breeding programmes for developing resistant varieties. Broad spectrum and durable resistance can be achieved by pyramiding of more than one resistance genes by marker assisted breeding programmes.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative Al technologies such as Large Language Models

(ChatGPT, COPILOT, etc) and text-to-image generators have been used during writing or editing of this manuscript.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Ausubel, F. M. (2005). Are innate immune signaling pathways in plants and animals conserved? *Nature Immunology*, 6, 973–979.
- Cao, J., Zhang, M., Xiao, J., Li, X., Yuan, M., & Wang, S. (2018). Dominant and recessive major R genes lead to different types of host cell death during resistance to *Xanthomonas oryzae* in rice. *Frontiers in Plant Science*, 9, 1711.
- Chen, X., Zuo, S., Schwessinger, B., Chern, M., Canlas, P. E., Ruan, D., Zhou, X., Wang, J., Daudi, A., Petzold, C. J., Heazlewood, J. L. (2014). An XA21-associated kinase (OsSERK2) regulates immunity mediated by the XA21 and XA3 immune receptors. *Molecular Plant*, 7(5), 874-892.
- Chu, Z., Yuan, M., Yao, J., Ge, X., Yuan, B., Xu, C., Li, X., Fu, B., Li, Z., Bennetzen, J. L., et al. (2006). Promoter mutations of an essential gene for pollen development result in disease resistance in rice. *Genes and Development*, 20, 1250-1255.
- da Cunha, L., McFall, A. J., & Mackey, D. (2006). Innate immunity in plants: A continuum of layered defenses. *Microbes and Infection*, 8(5), 1372-1381.
- de Wit, P. J. (2007). How plants recognize pathogens and defend themselves. *Cellular and Molecular Life Sciences*, 64, 2726-2732.
- Dodds, P. N., & Rathjen, J. P. (2010). Plant immunity: Towards an integrated view of plant-pathogen interactions. *Nature Reviews Genetics*, 11(8), 539-548.
- Gnanamanickam, S. S., Priyadarisini, V. B., Narayanan, N. N., Vasudevan, P., & Kavitha, S. (1999). An overview of bacterial blight disease of rice and strategies for its management. *Current Science*, 10, 1435-1444.
- Gopalakrishnan, S., Sharma, R. K., Anand Rajkumar, K., Joseph, M., Singh, V. P., Singh, A. K., Bhat, K. V., Singh, N. K., & Mohapatra, T. (2008). Integrating markerassisted background analysis with

foreground selection for identification of superior bacterial blight resistant recombinants in Basmati rice. *Plant Breeding*, 127(2), 131-139.

- Gu, K., Yang, B., Tian, D., Wu, L., Wang, D., Sreekala, C., Yang, F., Chu, Z., Wang, G. L., White, F. F., & Yin, Z. (2005). R gene expression induced by a type-III effector triggers disease resistance in rice. *Nature*, 435(7045), 1122-1125.
- Guvvala, L. D., Koradi, P., Shenoy, V., & Marella, L. S. (2013). Making an Indian traditional rice variety Mahsuri, bacterial blight resistant using marker-assisted selection. *Journal of Crop Science and Biotechnology*, 16, 111-121.
- Haney, C. H., Ausubel, F. M., & Urbach, J. M. (2014). Innate immunity in plants and animals: Differences and similarities. *The Biochemist*, 36(5), 40-45.
- Hu, K., Cao, J., Zhang, J., Xia, F., Ke, Y., Zhang, H., Xie, W., Liu, H., Cui, Y., Cao, Y., & Sun, X. (2017). Improvement of multiple agronomic traits by a disease resistance gene via cell wall reinforcement. *Nature Plants*, 3(3), 1-9.
- Iriti, M., & Faoro, F. (2007). Review of innate and specific immunity in plants and animals. *Mycopathologia*, 164, 57-64.
- Ji, Z., Wang, C., & Zhao, K. (2018). Rice routes of countering *Xanthomonas oryzae*. *International Journal of Molecular Sciences*, 19(10), 3008.
- Jiang, N., Yan, J., Liang, Y., Shi, Y., He, Z., Wu, Y., Zeng, Q., Liu, X., & Peng, J. (2020). Resistance genes and their interactions with bacterial blight/leaf streak pathogens (*Xanthomonas oryzae*) in rice (*Oryza sativa* L.)—An updated review. *Rice*, 13(1), 3.
- Jiang, Y., Cai, Z., Xie, W., Long, T., Yu, H., & Zhang, Q. (2012). Rice functional genomics research: Progress and implications for genetic crop improvement. Biotechnology Advances, 30, 1059-1070.
- Jones, J. D., & Dangl, J. L. (2006). The plant immune system. *Nature*, 444(7117), 323-329.
- Kesh, H., & Kaushik, P. (2020). Impact of marker assisted breeding for bacterial blight resistance in rice: A review, 151-165.
- Khush, G. S. (2005). What it will take to feed 5.0 billion rice consumers in 2030. *Plant Molecular Biology*, 59, 1–6.
- Khush, G. S., Bacalangco, E., & Ogawa, T. (1990). A new gene for resistance to

bacterial blight from *Oryza longistaminata*. *Rice Genetics News Letter*, 7, 121-122.

- Kumar, A., Kumar, R., Sengupta, D., Das, S. N., Pandey, M. K., Bohra, A., Sharma, N. K., Sinha, P., Sk, H., Ghazi, I. A., & Laha, G. S. (2020). Deployment of genetic and genomic tools toward gaining a better understanding of rice-*Xanthomonas oryzae* pv. *oryzae* interactions for development of durable bacterial blight resistant rice. *Frontiers in Plant Science*, 11, 1152.
- Kushalappa, A. C., Yogendra, K. N., & Karre, S. (2016). Plant innate immune response: Qualitative and quantitative resistance. *Critical Reviews in Plant Sciences*, 35(1), 38-55.
- Lee, S. W., Han, M., Park, C. J., Seo, Y. S., Bartley, L. E., & Jeon, J. S. (2011). The molecular mechanisms of rice resistance to the bacterial blight pathogen, *Xanthomonas oryzae* pathovar *oryzae*. In *Advances in Botanical Research* (Vol. 60, pp. 51-87). Academic Press.
- Lee, S. W., Han, S. W., Sririyanum, M., Park, C. J., Seo, Y. S., & Ronald, P. C. (2009). RETRACTED: A Type I–Secreted, Sulfated Peptide Triggers XA21-Mediated Innate Immunity. *Science*, 326(5954), 850-853.
- Liu, W., Liu, J., Triplett, L., Leach, J. E., & Wang, G. L. (2014). Novel insights into rice innate immunity against bacterial and fungal pathogens. *Annual Review of Phytopathology*, 52(1), 213-241.
- Liu, Y., Cao, Y., Zhang, Q., Li, X., & Wang, S. (2018). A cytosolic triosephosphate isomerase is a key component in XA3/XA26-mediated resistance. *Plant Physiology*, 178(2), 923-935.
- Mansfield, J., Genin, S., Magori, S., Citovsky, V., Sriariyanum, M., Ronald, P., Dow, M. A., Verdier, V., Beer, S. V., Machado, M. A., & Toth, I. A. (2012). Top 10 plant pathogenic bacteria in molecular plant pathology. *Molecular Plant Pathology*, 13(6), 614-629.
- Mew, T. W., Alvarez, A. M., Leach, J. E., & Swings, J. (1993). Focus on bacterial blight of rice. *Plant Disease*, 77, 5–12.
- Mizukami, T., & Wakimoto, S. (1969). Epidemiology and control of bacterial leaf blight of rice. *Annual Review of Phytopathology*, 7(1), 51-72.
- Mohapatra, S., Barik, S. R., Dash, P. K., Lenka, D., Pradhan, K. C., Raj, K. R., Mohanty, S. P., Mohanty, M. R., Sahoo, A., Jena, B. K., & Panda, A. K. (2023). Molecular breeding for incorporation of submergence tolerance and durable bacterial blight resistance into

the popular rice variety 'Ranidhan'. *Biomolecules*, 13(2), 198.

- Mundt, C. C. (2014). Durable resistance: A key to sustainable management of pathogens and pests. *Infection, Genetics and Evolution,* 27, 446-455.
- Muthamilarasan, M., & Prasad, M. (2013). Plant innate immunity: An updated insight into defense mechanism. *Journal of Biosciences*, 38, 433-449.
- Niño-Liu, D. O., Ronald, P. C., & Bogdanove, A. J. (2006). Xanthomonas oryzae pathovars: Model pathogens of a model crop. Molecular Plant Pathology, 7(5), 303-324.
- Noda, T., & Kaku, H. (1999). Growth of *Xanthomonas oryzae* pv. *oryzae* in planta and in guttation fluid of rice. *Japanese Journal of Phytopathology*, 65(1), 9-14.
- Nürnberger, T., Brunner, F., Kemmerling, B., & Piater, L. (2004). Innate immunity in plants and animals: Striking similarities and obvious differences. *Immunological Reviews*, 198(1), 249-266.
- Ou, S. H. (1985). *Rice diseases*. Commonwealth Mycological Institute.
- Pradhan, S. K., Nayak, D. K., Mohanty, S., Behera, L., Barik, S. R., Pandit, E., Lenka, S., & Anandan, A. (2015). Pyramiding of three bacterial blight resistance genes for broad-spectrum resistance in deepwater rice variety, Jalmagna. *Rice*, 8, 1-4.
- Rajpurohit, D., Kumar, R., Kumar, M., Paul, P., Awasthi, A., Osman Basha, P., Puri, A., Jhang, T., Singh, K., & Dhaliwal, H. S. (2011). Pyramiding of two bacterial blight resistance and a semidwarfing gene in Type 3 Basmati using marker-assisted selection. *Euphytica*, 178, 111-126.
- Sattari, A., Fakheri, B., Noroozi, M., & Moazami, K. (2014). Leaf blight resistance in rice: A review of breeding and biotechnology. *International Journal of Farming and Allied Sciences*, 3(8), 895-902.
- Singh, S., Sidhu, J. S., Huang, N., Vikal, Y., Li, Z., Brar, D. S., Dhaliwal, H. S., & Khush, G. S. (2001). Pyramiding three bacterial blight resistance genes (xa5, xa13, and Xa21) using marker-assisted selection into indica rice cultivar PR106. *Theoretical and Applied Genetics*, 102, 1011-1015.
- Song, W. Y., Wang, G. L., Chen, L. L., Kim, H. S., Pi, L. Y., Holsten, T., Gardner, J., Wang, B., Zhai, W. X., Zhu, L. H., & Fauquet, C. (1995). A receptor kinase-like protein encoded by the rice disease resistance gene, Xa21. *Science*, 270(5243), 1804-1806.

- Srinivasan, M. C., Thirumalachar, M. J., & Patel, M. K. (1959). Bacterial blight disease of rice. *Current Science*, 28, 469-470.
- Sundaram, R. M., Chatterjee, S., Oliva, R., Laha,
 G. S., Cruz, C. V., Leach, J. E., & Sonti, R.
 V. (2014). Update on bacterial blight of rice: Fourth international conference on bacterial blight. *Rice*, 7, 1-3.
- Sundaram, R. M., Vishnupriya, M. R., Biradar, S. K., Laha, G. S., Reddy, G. A., Rani, N. S., Sarma, N. P., & Sonti, R. V. (2008).
 Marker-assisted introgression of bacterial blight resistance in Samba Mahsuri, an elite indica rice variety. *Euphytica*, 160, 411-422.
- Verdier, V., Triplett, L. R., Hummel, A. W., Corral, R., Cernadas, R. A., Schmidt, C. L., Bogdanove, A. J., & Leach, J. E. (2012). Transcription activator-like (TAL) effectors targeting Os SWEET genes enhance virulence on diverse rice (*Oryza sativa*) varieties when expressed individually in a TAL effector-deficient strain of *Xanthomonas oryzae*. *New Phytologist*, 196(4), 1197-1207.
- White, F. F., & Yang, B. (2009). Host and pathogen factors controlling the rice-*Xanthomonas oryzae* interaction. *Plant Physiology*, 150(4), 1677-1686.
- Yang, B., Sugio, A., & White, F. F. (2006). Os8N3 is a host disease susceptibility gene for bacterial blight of rice. *Proceedings of the National Academy of Sciences of the United States of America*, 103, 10503-10508.
- Yoshimura, S., Yamanouchi, U., Katayose, Y., Toki, S., Wang, Z. X., Kono, I., Kurata, N., Yano, M., Iwata, N., & Sasaki, T. (1998). Expression of Xa1, a bacterial blightresistance gene in rice, is induced by bacterial inoculation. *Proceedings of the National Academy of Sciences*, 95(4), 1663-1668.
- Yoshimura, S., Yamanouchi, U., Katayose, Y., Toki, S., Wang, Z. X., Kono, I., Kurata, N., Yano, M., Iwata, N., & Sasaki, T. (1998). Expression of Xa1, a bacterial blightresistance gene in rice, is induced by bacterial inoculation. *Proceedings of the National Academy of Sciences*, 95(4), 1663-1668.
- Yuan, M., Chu, Z., Li, X., Xu, C., & Wang, S. (2010). The bacterial pathogen *Xanthomonas oryzae* overcomes rice defenses by regulating host copper redistribution. *Plant Cell*, 22, 3164-3176.

Sushitha and Sajeena; J. Adv. Biol. Biotechnol., vol. 27, no. 11, pp. 603-613, 2024; Article no. JABB. 125806

Zhang, H., & Wang, S. (2013). Rice versus Xanthomonas oryzae pv. oryzae: A unique pathosystem. *Current Opinion in Plant Biology*, 16(2), 188-195.

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of the publisher and/or the editor(s). This publisher and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.

© Copyright (2024): Author(s). The licensee is the journal publisher. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history: The peer review history for this paper can be accessed here: https://www.sdiarticle5.com/review-history/125806