A New Recessive Gene Conferring Resistance Against Rice Blast

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Abstract

Background: Rice blast (causative pathogen Magnaporthe oryzae) represents a major biotic constraint over rice production. While numerous genes for resistance have been found in both japonica and indica germplasm, as yet the diversity harbored by aus germplasm has not been widely exploited.

Results: The blast resistance present in the aus type cultivar AS20-1 was shown, via an analysis of segregation in the F2 generation bred from a cross with the highly blast susceptible cultivar Aichi Asahi, to be due to the action of a single recessive gene, denoted pi66(t). The presence of pi66(t) gave an intermediate level control to plants infected with the blast pathogen isolate EHL0635. A bulked segregant analysis indicated that four microsatellite loci (SSRs) mapping to chromosome 3 were probably linked to pi66(t). Localized mapping using chromosome 3-based SSRs and Indels defined a genetic window for pi66(t), flanked by the markers F04-j2 and M19-i12, which physically equals to 27.7 and 49.0 kb, respectively, in the reference genomes of cultivars Nipponbare and 93–11. This physical interval does not harbor any major gene currently associated with disease resistance.

Conclusion: pi66(t) is one of just three recessive genes controlling rice blast, and is the first major gene for resistance to be mapped to chromosome 3.

Keywords: Oryza sativa, Magnaporthe oryzae, aus rice cultivar, Recessive resistance gene

Background

Rice, a crop which feeds half of the world’s population, has been cultivated for at least 8,000 years (Khush 1997; The 3,000 rice genome project 2014; Travis et al. 2015). Five distinct groups of rice germplasm have long been recognized: they are referred to as indica, aus, basmati/sadri, tropical japonica and temperate japonica (The 3,000 Rice Genomes Project 2014; Travis et al. 2015). The aus group has developed in the north-eastern region of the Indian sub-continent, where both the climate and the growing environment are highly variable (Mahender et al. 2012; Travis et al. 2015). In recent years, aus germplasm has grown in importance as a source of genes for rice improvement, especially in the context of breeding for resistance/tolerance to abiotic and biotic stress (Travis et al. 2015 and references therein).

Rice blast (causative pathogen Magnaporthe oryzae) is a major constraint over rice production, inducing grain yield losses of up to 90% (He et al. 2012; Singh et al. 2015). Although breeders have so far been able to rely on a number of sources of genetic resistance, the pathogen is adept at evolving new races, with the result that monogenic resistances typically break down quite rapidly (Wu et al. 2014; Singh et al. 2015; Zhang et al. 2015). To date, some one hundred rice blast resistance (Pi) genes have been identified, many of which have been shown to map within a cluster or even in form of a tandem array; they are dispersed on eleven of the twelve rice chromosomes (Sharma et al. 2012; Singh et al. 2015; Tanwaeer et al. 2015 and references therein). All but two of the Pi genes are functionally dominant (Fukuoka et al. 2009; He et al. 2012), and about 30 have been isolated: their products mostly belong to the large group of nucleotide-binding site (NBS)-leucine-rich repeat (LRR) proteins. The two exceptions are Pid-2 and pi21 (Chen et al. 2006b; Fukuoka et al. 2009; Liu et al. 2011). Here, a third recessive gene, denoted pi66(t), has been identified...
in the *aus* cultivar (cv.) AS20-1, and its genomic position has been defined.

**Results**

**Resistance Reaction and Spectrum**

Numerous differential reactions were identified among the four cvs in the five *Mo* populations, suggesting that the *Pi* gene(s) carried by the donor cv. AS20-1 could be distinguished from the other *Pi* genes with these reactions (Table 1 and Additional file 1: Table S1). Intermediate and even lower resistance frequencies were evaluated among the four cvs in the five *M. oryzae* populations, indicating that all the four *Pi* genes should be incorporated with other *Pi* genes to stand the higher level of resistance in a given cultivar, if it will be released in the five *M. oryzae* populations.

**Resistance Inheritance**

When challenged by the blast isolate EHL0635, cv. AS20-1 was scored as moderately resistant (MR), cv. Aichi Asahi as susceptible (S) and the cv. AS20-1 x cv. Aichi Asahi F₁ as highly susceptible (HS) (Fig. 1). The qPCR-based assay confirmed that the hybrid was more susceptible than cv. Aichi Asahi. The F₂ progeny segregated as 101 R, 282 MR, 254 MS and 883 S, fitting a monogenic 1R:3S ratio when the R/MR and MS/S classes were combined ($\chi^2 = 0.02; P > 0.80$; Table 2). Together, these results indicated that the blast resistance expressed by cv. AS20-1 relied on homozygosity for the recessive allele of a single gene.

**Gene Locus**

BSA analysis revealed that four SSR markers (RM487, RM16, RM55, and RM168) on rice chromosome 3 were candidate markers linking to the target *Pi* gene, exclusively, in the F₂ population. The first round of linkage analysis with 750 viable F₂ plants revealed that there were 64 and 37 recombinants, respectively, at RM487 and RM16 loci on the centromere side, 35 and 22 distinct recombinants, respectively, at RM168 and RM55 loci on the telomere side, indicating that the four candidate markers were indeed linkage markers with the target *Pi* gene (Fig. 2a). Because no major *Pi* gene had been previously identified in this region, the novel *Pi* gene in AS20-1 was designated as *pi66(t)*.

Additional nine polymorphic SSR markers developed in the region defined by the flanking markers RM16 and RM55 were subjected to the second round of linkage analysis (Additional file 2: Table S2). The results showed that there were 31 to 22 recombinants detected among the seven marker loci [B07 (31), G02 (30), H15 (30), N03 (30), P23 (30), N11 (29), L18 (25), M23 (22)] on the centromere side, and only 8 distinct recombinants at RM135 locus on the telomere side (Fig. 2a). A total of 14 additional Indel markers developed in the narrower region flanked by markers M23 and RM135 were subjected to the third round of linkage analysis (Additional file 2: Table S2). The results showed that there were 15 to 2 recombinants detected among the six marker loci [D21 (15), E06 (9), I20 (7), I24 (2), F04 (2), F04-j2 (2)] on the centromere side, and 7 to 1 recombinant(s) detected among the eight marker loci [G23 (7), E01 (4), M19 (3), M19-4 (3), M19-3 (2), M19-2 (2), M19-1 (1), M19-i12 (1)] on the telomere side (Fig. 2a). The target locus, *pi66(t)*, was closely flanked by F04-j2 and M19-i12, which equals to 27.7 and 49.0 kb, respectively, in the reference genomes of cvs Nipponbare and 93–11 (Fig. 2a).

**Candidate Genes**

The *pi66(t)* region was represented by the two cv. Nipponbare overlapping BACs OSJNBB0009F04 and OSJN-Ba0092M19 (Fig. 2b). The number of genes present within this region was six in cv. Nipponbare and 14 in cv. 93–11 (Additional file 3: Table S3). Genome comparison and presence/absence (P/A) analyses revealed that there were three substantial Indel events that resulted in six genome-specific genes in the region. That is, both *pi66-2j* and *pi66-3j* in Indel I present in two genomes of cvs Nipponbare and AS20-1; *pi66-1i-2* (a duplication of *pi66-1i-1*) in Indel II, *pi66-5i* and *pi66-6i* in Indel III in that of both cvs 93–11 and AS20-1; and *pi66-2i* in Indel II in that of cv 93–11, only (Fig. 2c, Additional file 4: Figure S1).

![Table 1](image)

| Mo population | Selected isolate | Specific reactions selected from the five Mo populations* | Resistance frequencies in the five Mo populations (%)b |
|---------------|------------------|------------------------------------------------------|-----------------------------------------------------|
|               |                  | AS20-1 Aichi Asahi Kasalath IRBLta2-Pi                | AS20-1 Aichi Asahi Kasalath IRBLta2-Pi               |
| Guangdong     | CHL3417          | R          | R          | S          | 45.0 | 45.0 | 48.3 | 39.7 |
| Guangxi       | EHL1622          | S          | S          | R          | 25.0 | 26.7 | 36.7 | 38.3 |
| Yunnan        | EHL0210          | MS         | S          | MS         | 36.7 | 36.7 | 53.3 | 73.3 |
| Sichuan       | CHL892           | MR         | S          | S          | 20.0 | 20.0 | 48.3 | 56.7 |
| Heilongjiang  | EHL1379          | S          | S          | R          | 6.7  | 1.7  | 63.3 | 56.7 |

*a* R resistant, S susceptible, MS moderately susceptible, MR moderately resistant  
*b* Resistance frequencies were based on 60 isolates except for Kasalath and IRBLta2-Pi in the Guangdong population, in which only 58 isolates were tested.
Notably, there were six transposon-like genes (pi66-1j, -2j, -3j, 1i-1, 1i-2, -6i), of which both pi66-2j and -3j were scattered across the entire genomes except for the target region of cv. 93–11, thereby ruling out for P/A analysis (Additional file 4: Figure S1; Additional file 3: Table S3). Furthermore, there were three chimeric genes in both 93–11 and AS20-1 genomes (Fig. 2c and Additional file 4: Figure S1). By excluding six transposon-like genes, there were three most possible candidates (pi66-5a, -6a, -7a) for pi66(t) (Fig. 2c; Additional file 3: Table S3).

**Discussion**

Chinese rice breeders have to date largely ignored aus germplasm, even though it has acquired a growing reputation for harboring genes for resistance/tolerance to abiotic and biotic stress (Travis et al. 2015). Rather, efforts to improve indica have concentrated on materials developed in SE Asia, while those directed at japonica have relied on germplasm from Japan (Wu et al. 1991). In addition to pi66(t), aus germplasm has also yielded both Pi16 and an allele of Pik (Pan et al. 1999). More recently, nine already recognized Pi genes have been identified as present in materials originating in NE and E India (Imam et al. 2014), which is the center of origin of aus germplasm. Notably, the donor of pi66(t) also harbors a gene conferring resistance against the brown plant hopper; this gene also lies on chromosome 3, but at some distance from pi66(t) (Chen et al. 2006a). Such works clearly indicated that aus cvs are valuable and promising genetic resources for withstanding biotic pressures including rice blast disease, and will greatly enlarge the gene pool for rice breeders.

Plant disease resistance genes have been classified into two types, the most frequent of which encode an NBS-LRR protein. Non-NBS-LRR genes encode a wide diversity of products (Chauhan et al. 2015; Olukolu et al. 2016), tend to confer partial (rather than complete) resistance and are typically more durable than the NBS-LRR type genes. The most well documented non-NBS-LRR type is barley mlo, a gene which encodes a G protein-coupled receptor residing in the plasma membrane (Kime et al. 2002); the gene confers durable resistance to a broad spectrum of powdery mildew races (Acevedo-Garcia et al. 2014). A second example is the wheat gene Lr34, which encodes an ATP-binding cassette transporter; its product protects against five distinct foliar fungal pathogens (Krattinger et al. 2009; Chauhan et al. 2015).

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**Table 2** Segregation for resistance in the F2 population bred from the cross cv. AS20-1 x cv. Aichi Asahi, following inoculation with the M. oryzae isolate EHL0635

| Parents/F2 plants | No. of plants | Segregation | χ² | P |
|------------------|---------------|-------------|----|---|
|                   | R  | MR | MS | S | Total |
| AS20-1            | 9  | 19 | 1  | 0 | 29    |
| Aichi Asahi       | 0  | 0  | 0  | 3 | 3     |
| F2 population     | 101| 282| 254| 83| 1520  |

*R resistant, S susceptible, MS moderately susceptible, MR moderately resistant

*na not applicable

Chi-square test using the Yates correction comparing resistance [R + MR] with susceptibility [MS + S]
The maize gene *ZmWAK* encodes a plasma membrane-related receptor-like kinase; its presence has been correlated with a reduction in the incidence of head smut disease (Zuo et al. 2015). Finally, the rice gene *xa5* encodes a small subunit of the transcription factor IIA (TFIIA); this gene confers resistance against bacterial blight (Iyer-Pascuzzi 2004). Before the identification of *pi66*(t), all but two of the *Pi* genes characterized to date act as dominant alleles. The exceptions are *pi21* and *pi55*. The former gene encodes a proline-rich protein harboring a probable heavy metal-binding domain and some predicted protein-protein interaction motifs; the resistant allele differs from the wild type dominant one by two deletions affecting the latter motifs, and which are thought to be responsible for the allele’s determination of non-race-specific resistance (Fukuoka et al. 2009). One of candidate genes for *pi55* encodes a protein rather similar to that encoded by *pi21*. Although a substantial number of major *Pi* genes have been intra-chromosomally mapped, *pi66*(t) is the first to be located on chromosome 3. Two quantitative trait loci mapping to this chromosome (*Os03g0122000* and *Os03g0120400*) have been associated with blast resistance (Wang et al. 2014), but both lie outside the critical RM16-RM55 interval. The *pi66* identified in the current study that is the third recessive *Pi* gene located on the virgin land, where no any known *Pi* protein (domain) is identifiable (Additional file 3: Table S3). It is noteworthy that the *bph19* derived from the donor cv. AS20-1 was also recognized as non-NBS-LRR resistance gene (Chen et al. 2006a). It has been argued that durable and broad-spectrum resistance may be more readily achieved by deploying non-NBS-LRR genes, perhaps in combination with NBS-LRR ones, than by attempting to stack genes which each (at least for some time) confer immunity (Fukuoka et al. 2009; Acevedo-Garcia et al. 2014; Chauhan et al. 2015; Zuo et al. 2015). This hypothesis can only be tested by exploiting genes such as *pi66*(t) in a rice breeding program.

**Conclusions**

This research has confirmed that novel resistance genes against blast can be recovered from *aus* germplasm. The gene *pi66*(t) identified here is the third recessive *Pi* gene to be identified, and is also the first major *Pi* gene to be located on chromosome 3.

**Methods**

**Phenotyping**

The *pi66*(t) donor cv. AS20-1, along with the *Pia* carrier cv. Aichi Asahi, the *Pi36* carrier cv. Kasalath and the *Pita*-2 carrier cv. IRBLta2-Pi were challenged with 60 *M. oryzae* isolates collected from each of Guangdong (GD), Guangxi...
Chromosomal Mapping
The donor cv. AS20-1 was crossed with the highly susceptible cv. Aichi Asahi, and their F₂ progenies were screened for reaction to inoculation with M. oryzae isolate EHL0635. The F₂ population showing monogenic segregation was regarded as the mapping population, thereby subjecting to the bulked-segregant assay (BSA) for quickly mapping chromosomal region involving the target gene. Genomic DNAs of the F₂ plants as well as the parental plants were extracted from frozen leaves using the CTAB method. Two contrast bulks that were constructed by pooling equimolar amounts of DNAs from 10 resistant or susceptible F₂ plants. The two bulks, along with both parental DNAs, were then assayed with a set of 180 simple sequence repeat (SSR) markers (Temnykh et al. 2000, 2001), selected to span the full rice genome, following the methods given by He et al. (2012).

Gene Mapping
Genomic map of target gene were established through three rounds of linkage analysis using genomic position-ready molecular markers (He et al. 2012). The first round was carried out with candidate markers defined by BSA for screening recombinants on both sides of the target locus. The second round was carried out with additional SSR markers in the target region flanked by the closest markers derived from the first round of linkage analysis, which were developed on the basis of reference sequence of cv. Nipponbare, except for RM135 that was adopted from the rice SSR marker maps (Temnykh et al. 2000, 2001). The third round was carried out in the recombinant progeny with insertion/deletion (Indel) markers those were developed de novo based on differential sequences between the two reference sequences of japonica cv. Nipponbare and indica cv. 93–11. Linkage marker search and prime designation were performed in the way essentially same way as previously described (Liu et al. 2005; Zeng et al. 2011; He et al. 2012). Genomic map of the target locus was constructed on the basis of both reference sequences.

Candidate Gene Indentification
Candidates for pi66(t) were predicted based on gene annotations provided by BLASTN (www.ncbi.nlm.nih.gov/BLAST), RiceGAAS (ricegaas.dna.affrc.go.jp) and FGENSH (www.softberry.com) software. The two reference sequences proved to be rather diverse in the target region, so candidates that encode proteins with over 200 aa was validated by PCR-based presence/absence (P/A) test against the four DNAs of cvs AS20-1, Aichi Asahi, 93–11, and Nipponbare, following Zhai et al. (2011).

Additional Files

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Competing Interests
The authors declare that they have no competing interests.

Authors’ Contributions
Project conception (QP). Resistance phenotyping (QP, LW). Mapping population construction (QP, LW). Linkage analysis (2L, QP). Candidate gene analysis (2L, QP). Manuscript preparation (QP). All authors read and approved the final manuscript.

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