

Figure S1. Disease resistance to *Xanthomonas oryzae* pv. *oryzae* (T7174, race I) in P_{OsUbi7} -*BSR1* T1 lines. Top leaves (L6-L8) of P_{OsUbi7} -*BSR1* and WT lines were inoculated with *X. oryzae* pv. *oryzae* (T7174). The arrowhead indicates the point of inoculation.

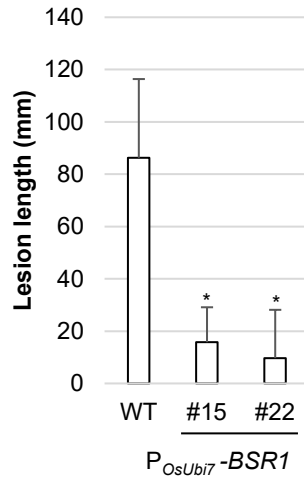


Figure S2. Disease resistance to *Xanthomonas oryzae* pv. *oryzae* (T7133, race III) in $P_{OsUbi7}\text{-}BSR1$ T1 lines. Top leaves (L6-L8) of $P_{OsUbi7}\text{-}BSR1$ and WT plants were inoculated with *X. oryzae* pv. *oryzae* (T7133). Lesion lengths in $P_{OsUbi7}\text{-}BSR1$ lines were significantly lower than those in wild-type (WT) plants (* $P < 0.05$ by Dunnett's test). Values are mean \pm SD (n = 6-18).

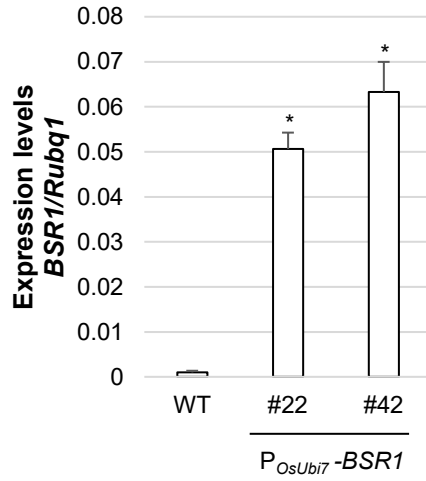


Figure S3. *BSR1* expression levels in P_{OsUbi7}-*BSR1* T4 lines. *BSR1* expression levels in P_{OsUbi7}-*BSR1* lines were significantly higher than those in wild-type (WT) plants (* $P < 0.05$ by Dunnett's test). Values are mean \pm SD (n = 3).

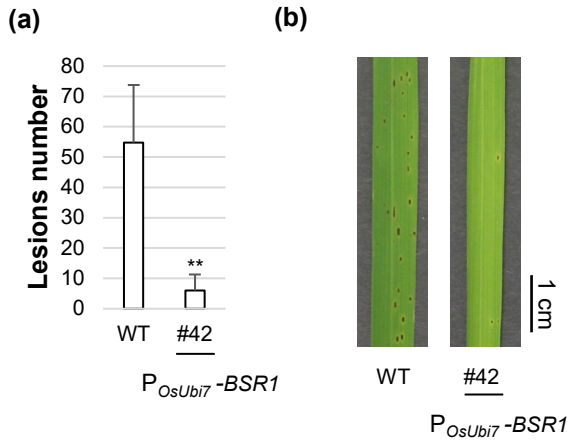


Figure S4. Disease resistance to *Cochliobolus miyabeanus* in P_{OsUbi7} -BSR1 T4 line. (a) Lesion numbers on *C. miyabeanus* infected T4 leaves in P_{OsUbi7} -BSR1 and WT lines 4 d after inoculation. The inoculum concentration was 5×10^4 conidia/ ml. Lesion numbers in P_{OsUbi7} -BSR1 plants were significantly lower than in WT plants 4 d after inoculation (** $P < 0.01$ by t-test). Values are mean \pm SD (n = 6-7). (b) Photographs of leaves infected with *C. miyabeanus* in P_{OsUbi7} -BSR1 and WT lines 4 d after inoculation.

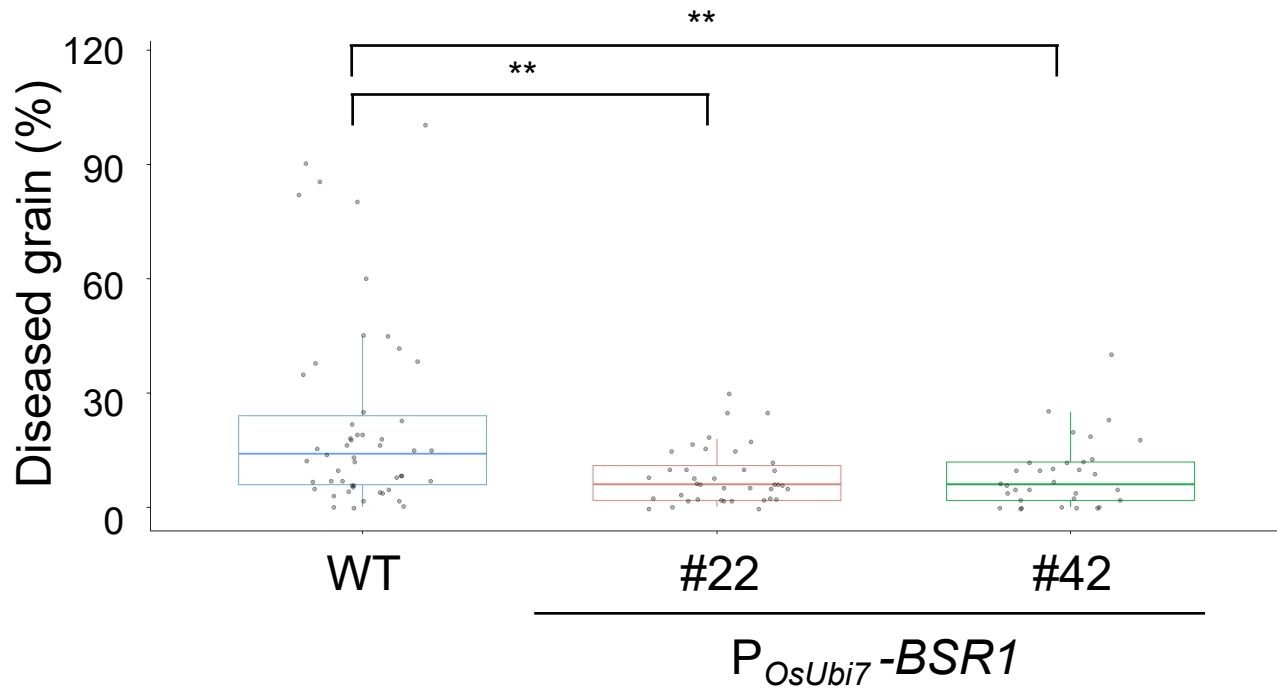


Figure S5. Disease resistance to panicle blast caused by *Pyricularia oryzae* (isolate Kyu89-246) in the descendants of the $P_{OsUbi7-BSR1}$ #22 and #42 lines. Kruskal-Wallis test with Steel's post-hoc test was used to analyze significant differences between these group (** $P < 0.01$).

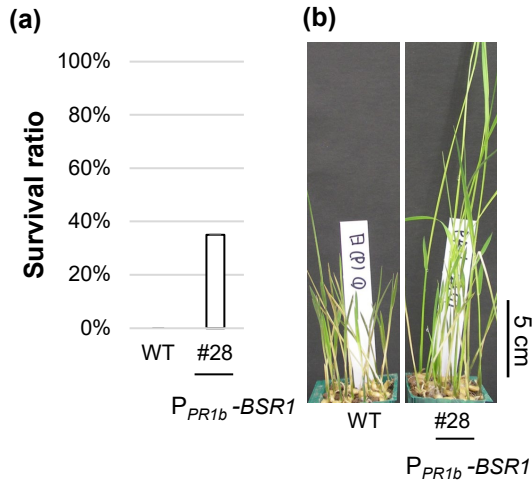


Figure S6. Disease resistance to *Burkholderia glumae* in P_{PR1b}-BSR1 T2 lines. (a) Pre-germinated T2 seeds of P_{PR1b}-BSR1 and wild-type (WT) lines were inoculated with *B. glumae*. The inoculum concentration was OD₅₂₀ = 0.0004. Survival ratio was calculated 8 d after inoculation (n = 20). Tests were repeated thrice with similar results. (b) Photographs of *B. glumae* infected shoots in P_{PR1b}-BSR1 and WT lines 8 d after inoculation.

Method S1. Evaluation of panicle blast resistance

The young seedlings were transplanted into plastic pots, grown in the glasshouse room of the NARO, and used for inoculation within 10 d from the day when the neck of the panicle emerged. Spores of race 003.0, a blast fungus pathogenic to WT and P_{OsUbi7}-*BSRI* lines, were suspended in 0.01% Tween 20 and sprayed with 20 mL per pot at a concentration of 1.0×10^5 conidia/ mL; and the plants were kept in a dew chamber for 20 h at 24.5 °C and grown in a glasshouse. The proportions of diseased grains per panicle were examined approximately 3 weeks after inoculation [1, 2]. Statistical analyses were conducted using EZR (Saitama Medical Center, Jichi Medical University, Saitama, Japan), which is a graphical user interface for R [3]. The Kruskal–Wallis test with Steel’s post-hoc test was used to analyze significant differences between these group (** $P < 0.01$).

References

1. Hayashi, N.; Inoue, H.; Kato, T.; Funao, T.; Shiota, M.; Shimizu, T.; Kanamori, H.; Yamane, H.; Hayano-Saito, Y.; Matsumoto, T.; Yano, M.; Takatsuji, H., Durable panicle blast-resistance gene Pb1 encodes an atypical CC-NBS-LRR protein and was generated by acquiring a promoter through local genome duplication. *Plant J.* **2010**, 64, (3), 498-510.
2. Inoue, H.; Hayashi, N., The Panicle Blast Resistance Mechanism of in the Rice Cultivar Miyazaki-mochi is Independent from that of. *Jarq-Jpn Agr Res Q* **2019**, 53, (4), 289-293.
3. Kanda, Y., Investigation of the freely available easy-to-use software 'EZR' for medical statistics. *Bone Marrow Transplant.* **2013**, 48, (3), 452-458.