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Research Article

Genetic Performance of the Semidwarfing Allele \textit{sd1} Derived from a Japonica Rice Cultivar and Minimum Requirements to Detect Its Single-Nucleotide Polymorphism by MiSeq Whole-Genome Sequencing

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The influence of the semidwarfing gene \textit{sd1} derived from the rice cultivar Jukkoku (Jukkoku\textit{sd1}) and IR8 (IR8\textit{sd1}), which contributed to the Green Revolution, \textit{d60} from Hokuriku 100, as well as the combination of \textit{sd1} and \textit{d60} (Jukkoku\textit{sd1 plus d60} and IR8\textit{sd1 plus d60}), was investigated using isogenic lines raised by backcrossing with the cultivar Koshihikari. The isogenic lines carrying Jukkoku\textit{sd1}, IR8\textit{sd1}, \textit{d60}, Jukkoku\textit{sd1 plus d60}, and IR8\textit{sd1 plus d60} had considerably shorter culm lengths than Koshihikari by 19.2%, 22.8%, 26.0%, 45.1%, and 43.4%, respectively. The \textit{sd1 plus d60} lines showed additively reduced culms, indicating that the function of \textit{d60} was different from \textit{sd1}. In contrast to the culm reduction, Jukkoku\textit{sd1} showed productive merit with a panicle length of 2.5% greater than the origin. MiSeq next-generation sequencer was used to optimize a minimum scale to detect Jukkoku\textit{sd1} in practical breeding. Mapping with the reference genome of Nipponbare gained the average depths of Koshihikari Jukkoku\textit{sd1} and Koshihikari being 9.17 and 7.29, respectively. Comparing the vcf files of the entire genomes of Koshihikari Jukkoku\textit{sd1} and the virtual Koshihikari revealed a G to T SNP at position 38,382,746 in the \textit{sd1} locus on chromosome 1 of Koshihikari, causing a loss-of-function mutation of GA20-oxidase.

1. Introduction

Semidwarfism of rice improves phenotype (light-interception properties and harvest index), nitrogen responsiveness, and lodging resistance and is thus an important trait worldwide. The International Rice Research Institute produced a cross between Peta, a tall \textit{indica} variety (culm length, 150–180 cm), characterized by abundant, long hanging leaves, commonly grown in tropical Asia, and Dee-geo-woo-gen (DGWG), a Taiwanese indigenous semidwarf variety, to improve its lodging resistance and light-interception properties. The resulting semidwarf rice variety IR8 (culm length, 90–100 cm), developed in 1966, has dramatically improved rice yields and brought the Green Revolution to tropical Asia [1].

The semidwarf trait has also been introduced into rice cultivars grown in other countries. In Japan, Hoyoku, Shiranui, and Reihou, which are representative main cultivars grown in the Kyushu region, were developed using the indigenous semidwarf cultivar Jukkoku [2]. A series of cultivars, such as Akihikari and Niigata Wase, were developed in the 1970s in the Tohoku region using the semidwarf cultivar Reimei, which was induced by gamma-ray irradiation of Fujiminori [3]. In the USA, Calrose 76 (culm length, ~90 cm) was developed in 1976 also by gamma-ray irradiation of the \textit{japonica} variety Calrose (culm length, ~120 cm) [4–6]. In Korea, \textit{d47} derived from IR8 was also introduced into 'Longil' by japonica-indica hybridization [7].

Advances in genetic research have facilitated the identification of genes responsible for semidwarfism in rice. The incomplete recessive gene \textit{d47} is responsible for the semidwarfism of DGWG, the parent line of IR8 [8–10]. Later, the incomplete recessive gene \textit{sd1} in Calrose 76 was shown...
to be allelic to *d47* [4, 11, 12]. Moreover, allelic relationships of the semidwarf gene have been found in Taichung Native 1, derived from DGWG; Shiranui, derived from Jukkoku; and *d49* in the mutant cultivar Reimei [13, 14]. Finally, all these semidwarf cultivars carry alleles at the *sd1* locus, despite their different parentage [14–18]. The *sd1* alleles, on the long arm of chromosome 1, encode loss-of-function mutations in GA20-oxidase (*OsGA20ox2*), which regulates the synthesis of biologically active gibberellins (GAs), which catalyze three steps in the GA biosynthesis pathway [15–18]. Taken together, a single semidwarf gene, *sd1*, solely confers the semidwarf phenotype of cultivars commonly grown around the world due to its nondetrimental effects on grain yield [19].

This narrow genetic base of current semidwarf rice cultivars has led to reduced genetic diversity in rice [20–22]. Thus, it is necessary to identify a novel semidwarf gene alternative to *sd1* and to utilize it to extend genetic diversity in semidwarf cultivars. A novel dwarf gene, *d60*, which was found in the semidwarf mutant Hokuriku 100, developed by exposing Koshihikari to 20 kR of gamma radiation [23, 24], is thus of particular importance. Unlike *sd1*, which is inherited as a single recessive gene, *d60* complements the gametic lethal gene *gal*, which is carried by many rice varieties. Consequently, the cross between Hokuriku 100 and Koshihikari exhibits a unique genotype ratio of 4*D60*D60 : 4*D60*d60 : 1*d60*d60 [23–25].

Although many semidwarf genes allelic to *sd1* have been identified in different cultivars (DGWG, Jukkoku, Reimei, and Calrose 76), differences in their influences have not been elucidated. Thus, investigating the differences in ecological and phenotypic traits in relation to yield between *d60*- and *sd1*-carrying NILs, in relation to yield, to assess the utility of *sd1* and *d60* semidwarf genes and (b) sequence NILs by MiSeq to detect the differences between *Sd1*/Sd1* target sites.

### 2. Materials and Methods

#### 2.1. Plant Genetic Materials

As shown in Figure 1, the following rice cultivars were examined: Koshihikari, Koshihikari Jukkoku_ *sd1* (Koshihikari*+*//Koshihikari/Jukkoku _B6_F3_), Koshihikari IR8_ *sd1* (Koshihikari*+*//Koshihikari/ Kinuhikari B6_F3), Kinuhikari carrying *sd1* of IR8 [26], Koshihikari *d60* (Koshihikari*+*//Koshihikari/Hokuriku 100 B6_F3), Koshihikari Jukkoku_ *sd1* plus *d60* (Koshihikari Jukkoku_ *sd1*/Koshihikari *d60*), and Koshihikari IR8_ *sd1* plus *d60* (Koshihikari IR8_ *sd1*/Koshihikari *d60*). In each backcross experiment, short-culm BCn_F3 plants were backcrossed with Koshihikari as the recurrent female parent in each generation. Each semidwarf phenotype was fixed by the BCn_F3 generation. The percentage of genetic materials from Koshihikari in the isogenic semidwarf lines was calculated as follows: For Koshihikari carrying Jukkoku_ *sd1* (B6_F3), (1 – (1/2)6) × 100 = 98.4%. For B6_F3, (1 – (1/2)2) × 100 = 87.5%.

#### 2.2. Cultivation

Rice seeds were collected from stocks kept in a refrigerator. Seeds of each line were immersed in enough water to cover them. Water was changed every day for 7 days (May 2 to May 8) during germination stimulation. Seedlings were grown in special boxes (30 × 15 × 3 cm) for approximately 20 days in a greenhouse: two seeds were planted in each cell (2 × 2 × 3 cm) in the box on May 11 and 12 and watered twice a day (07:00 and 12:00). Seedlings were then individually transplanted into a paddy field (120 m²: 6.0 × 20.0 m) of the University Farm on June 8. The paddy field received 4.0 kg of basal fertilizer containing nitrogen, phosphorus, and potassium (weight ratio, nitrogen : phosphorus : potassium = 2.6 : 3.2 : 2.6). Except for the period of mid-season drainage (July 10 to July 17), the water level was maintained at 5–7 cm above ground to prevent seedlings from submersing. A herbicide (Joystar L floable).
was applied on June 20 to kill weeds growing uncontrollably and the water was then kept at a high enough level to cover the weeds for 1 week. Koshihikari, Koshihikari carrying Jukkoku sd1, Koshihikari carrying IR8 sd1, Koshihikari carrying d60, Koshihikari carrying d60 and Jukkoku sd1, and Koshihikari carrying d60 and IR8 sd1 were grown in 4 m² plots (2 × 2 m) (two instances per line). Basal fertilizer was applied to give 0.43 g/m² nitrogen, 0.53 g/m² phosphorus, and 0.53 g/m² potassium. After ripening, 10 plants per genotype were sampled twice to assess the following traits: panicle heading time, culm length, internode interval, and panicle length.

2.3. Investigation of Agronomic Traits. The time when the tip of the panicle first emerged from the flag leaf sheath was recorded as the heading time for all plants. Ten plants typical of each line were selected. The sampling procedure was performed twice. Sampled plants were air-dried and the following phenotypic traits were assessed or measured: (1) culm length (the length between the ground surface and the panicle end of the main culm), (2) internode length (the lengths between two neighboring internodes of the upper five internodes), and (3) panicle length (the length between the panicle base and the tip of the panicle). Differences in phenotypic traits between each line carrying a semidwarf gene (or genes) and Koshihikari were obtained using the following equation: percent difference = [(measurements of Koshihikari) – (measurements of each line)]/(measurements of Koshihikari) × 100.

2.4. Next-Generation Sequencing (NGS) Analysis. An advantage of genomics is the development of a next-generation sequencer that can decode DNA sequences at the gigal level. Development of the next-generation DNA sequencer was advanced under the Affordable Care Act aims to realize societal implementation of medical genomics [27, 28]. The desktop-type next-generation sequencer MiSeq has the ability to read 15 million DNA sequences in one run. Generally, whole-genome sequencing analysis required 30 times of the standard scale, the MiSeq treats only one rice sample per run, because the 15 million DNA amount is just enough to reconstruct a single rice genome. However, it is a big problem that the running cost is excessively high to use the MiSeq at the practical breeding to detect target genes. So, in this study, to achieve a minimum scale to detect target genes with a reasonable cost, we try to detect Jukkoku sd1 at only 5x rice genome coverage by using the MiSeq. The semidwarf gene sd1 (a loss-of-function mutation of the GA20-oxidase encoding gene) was transferred to Koshihikari by consecutive backcrosses to prepare a semidwarf Koshihikari named Kukkoku sd1. The Koshihikari Jukkoku sd1 backcross was used to detect single-nucleotide polymorphisms (SNPs) in Jukkoku-derived sd1 by NGS. Whole-genome analysis was conducted using Koshihikari Jukkoku sd1 and Koshihikari. Genomic DNA was extracted from each cultivar by the CTAB (hexadecyltrimethylammonium bromide) method. Genomic DNA was tagged and fragmented to an average length of 500 bp using the Nextera® transposome-based approach. After purification of the transposome with DNA Clean & Concentrator™-5 (Zymo Research, Irvine, CA, USA), adaptors for fixation on the flow cell were synthesized at both ends of each fragment using a polymerase chain reaction (PCR). Then, the DNA fragments were subjected to size selection using AMPure XP magnetic beads (Beckman Coulter, Inc., Brea, CA, USA). Finally, qualitative and quantitative measurements were performed using a Fragment Analyzer™ (Advanced Analytical Technologies, Inc., Ankeny, IA, USA) and a Qubit® 2.0 Fluorometer (Thermo Fisher Scientific, Waltham, MA, USA) to prepare a DNA library for NGS. Aiming to achieve 5× rice genome coverage, a MiSeq next-generation sequencer was used to simultaneously analyze five lines; namely, 4.5 ng of five DNA libraries was applied in each MiSeq run. Clusters then were formed on the flow cells by bridge-PCR and each pair-end of a 250 bp read was sequenced. Resulting sequenced reads were mapped using BWA software with the Nipponbare genome [29] as a reference. Then, SNPs and Indels were detected using SAMtools software (http://samtools.sourceforge.net/).

3. Results

3.1. Effects of Semidwarf Genes sd1 and d60 on Heading Time. The days to heading were compared (Figure 2). The earliness of varieties of Koshihikari and donors of semidwarf genes is as follows: Koshihikari, an early-medium maturing variety; Jukkoku, a medium-late maturing variety; Kinuhikari, an early-medium maturing variety; and Hokuriku 100, an early-medium maturing variety. The heading panicles were observed first in Koshihikari carrying IR8 sd1 (83 days to heading) and the latest heading panicles were observed in Koshihikari carrying Jukkoku sd1 or Jukkoku sd1 plus d60 (93 days to heading). However, the difference in the average number of days to heading was the largest between lines carrying IR8 sd1 (86.5 days) and those carrying d60 (90.5 days), but this 4-day difference was thought to be minor (Figure 2). Thus, it was concluded that the time required for maturing was comparable among lines and the differences
Table 1: Effects of the semidwarfing genes Jukkoku_sd1, IR8_sd1, and d60 and their combinations on panicle length, culm length, and internode interval.

| Gene               | Panicle length (cm) | Culm length (cm) | 1st internode interval (cm) | 2nd internode interval (cm) | 3rd internode interval (cm) | 4th internode interval (cm) | 5th internode interval (cm) |
|--------------------|---------------------|------------------|---------------------------|---------------------------|---------------------------|---------------------------|---------------------------|
| Koshihikari        | 16.2                | 88.8             | 39.7                      | 20.7                      | 17.8                      | 11.6                      | 3.1                       |
| Jukkoku_sd1        | 16.6 (+2.5%)        | 71.8 (-19.2%)    | 30.2                      | 15.0                      | 13.0                      | 8.8                       | 2.5                       |
| IR8_sd1            | 15.8 (-2.4%)        | 68.5 (-22.8%)    | 33.2                      | 14.2                      | 11.2                      | 6.6                       | 1.7                       |
| d60                | 15.7 (-3.0%)        | 65.7 (-26.1%)    | 30.1                      | 14.0                      | 12.9                      | 7.6                       | 1.7                       |
| Jukkoku_sd1 + d60  | 15.8 (-2.2%)        | 48.6 (-45.3%)    | 26.7                      | 10.7                      | 7.9                       | 3.1                       | 0.5                       |
| IR8_sd1 + d60      | 15.7 (-3.3%)        | 50.2 (-43.5%)    | 30.1                      | 11.2                      | 6.6                       | 1.9                       | 0.3                       |

Figure 3: Effects of the semidwarfing genes Jukkoku_sd1, IR8_sd1, and d60 and their combinations on internode constitution.

Figure 4: Effects of the semidwarfing genes Jukkoku_sd1, IR8_sd1, and d60 and their combinations on panicle length.

in traits, such as panicle length, were attributed to genetic factors.

3.2. Effects of Semidwarf Genes sd1 and d60 on Traits Related to Plant Type. The mean values of the individual traits in plants of each genotype were calculated and the percent differences relative to the corresponding mean values of Koshihikari were compared. Introduction of a semidwarf gene (or genes) resulted in a reduction in culm length. The mean culm length of Koshihikari was 88.8 cm, while those of lines carrying Jukkoku_sd1, IR8_sd1, d60, Jukkoku_sd1 plus d60, and IR8_sd1 plus d60 were 71.8, 68.5, 65.7, 48.6, and 50.2 cm, respectively (Table 1 and Figure 3). Similarly, internode intervals were also reduced by the introduction of at least one semidwarf gene. While reductions in internode intervals were relatively uniform in lines carrying Jukkoku_sd1 (23.9%, 27.5%, 27.0%, 23.3%, and 19.4%; percent difference, from upper to lower internode intervals), the reduction in length between the third and fourth and between the fourth and fifth internodes was markedly large in lines carrying IR8_sd1 (16.4%, 31.5%, 37.0%, 42.6%, and 45.0%; percent difference, from upper to lower internode intervals). Reductions in internode intervals were relatively uniform in lines carrying d60 (24.2%, 32.6%, 27.4%, 34.5%, and 45.7%; percent difference, from upper to lower internode intervals), while percent differences were larger than those observed in lines carrying Jukkoku_sd1. In the sd1 plus d60 lines, marked reductions were observed in length between neighboring internodes, probably attributed to the additive effect of sd1 and d60 (Table 1 and Figure 3). The interval between the fourth and fifth internodes often disappeared in plants carrying IR8_sd1 and in sd1 plus d60 lines, which showed a marked reduction in length between the lower internodes.

In contrast, as shown in Figure 4, the effect of a semidwarf gene (or genes) on panicle length (−3.0 to +2.5%), which
was smaller than that for internode intervals (~16 to ~92%), was negligible in practical agriculture. Particularly, the mean length of panicles in the line carrying Jukkoku_sd1 was 16.2 cm, which was solely 2.5% longer than that of Koshihikari. The panicle length of the line carrying Jukkoku_sd1 was longer than that of the original cultivar Koshihikari (Table 1 and Figure 4). However, the reduction in panicle length was considerably less than that in culm length (Table 1 and Figure 3; a 22.8% decrease at o t a l r e a d y 2.5% in Koshihikari carrying Jukkoku_sd1, while a 26.1% decrease in semidwarf lines had shorter culm lengths than Koshihikari, with the original cultivar Koshihikari (Table 1 and Figure 4). In this study, the G to T SNP of the defective GA20-oxidase gene was localized at position 38,382,746 from the end of the short arm of chromosome 1 of Koshihikari. Using this scale of the NGS approach with a coverage of only 5x, the targeted SNP in Jukkoku_sd1 was successfully identified with three reads, with a cost reduced to 1/5 that of the ordinal whole-genome sequencing with a coverage of 30x.

### 3.3. NGS Analysis to Detect Jukkoku-Derived sd1 with a Minimum Scale.

Using the MiSeq sequencer, we obtained a total read length of 2.79 × 10^6 bp from the total read number of 9.53 × 10^6 in Koshihikari Jukkoku_sd1, while a total read length of 1.92 × 10^6 bp was obtained from a total read number of 6.13 × 10^6 in Koshihikari. By mapping the read sequences obtained by NGS using Nihonbare genomic DNA as a reference, sequence coverage rates of 93.5% and 88.5% were attained for Koshihikari Jukkoku_sd1 and Koshihikari, respectively, while average depths were 9.17 and 7.29, respectively. Furthermore, vcf files of the entire genomes were prepared and the whole-genome sequence of Koshihikari Jukkoku_sd1 was compared with that of the virtual Koshihikari genome. As a result, three reads were obtained, including Jukkoku_sd1 from Koshihikari Jukkoku_sd1, while three reads of Sd1 were obtained from Koshihikari. The Sd1/sd1 locus (Os01t0883800-01) was localized at positions 38,382,385–38,385,469 from the end of the short arm of chromosome 1 in the Koshihikari genome. The difference observed between the Koshihikari_Sd1/Jukkoku_sd1 alleles was only one SNP from G to T in exon 1 of the GA20-oxidase gene (Figure 5), as reported by Sasaki et al. [17]. In this study, the G to T SNP of the defective GA20-oxidase gene was localized at position 38,382,746 from the end of the short arm of chromosome 1 of Koshihikari. Using this scale of the NGS approach with a coverage of only 5x, the targeted SNP in Jukkoku_sd1 was successfully identified with three reads, with a cost reduced to 1/5 that of the ordinal whole-genome sequencing with a coverage of 30x.

### 4. Discussion

Environmental degradation caused by global warming, postearthquake salt damage, and radioactive contamination and globalization of agricultural markets due to the Trans-Pacific Partnership are serious issues that call for the innovation of new cultivars to overcome the shortcomings of current crops. The results of this study showed that all tested semidwarf lines had shorter culm lengths than Koshihikari, indicating improved lodging resistance. Furthermore, the leaves were straighter (pointing upwards) in the semidwarf lines than in Koshihikari (Figure 1), indicating improved light-interception properties attributed to the introduction of semidwarf gene(s). Panicle length was solely longer by 2.5% in Koshihikari carrying Jukkoku_sd1 and shorter in lines carrying IR8_sd1 by 2.4% or d60 by 3.0%, as compared with the original cultivar Koshihikari (Table 1 and Figure 4). However, the reduction in panicle length was considerably less than that in culm length (Table 1 and Figure 3; a 22.8% decrease in lines carrying IR8_sd1 versus a 26.1% decrease in lines carrying d60), suggesting that the negative effects
of the semidwarf genes sd1 and d60 on panicle length were negligible (Table 1 and Figure 4). Ogi et al. [30] and Murai et al. [31] reported that an isogenic line carrying sd1 was derived from DGWG, which was constructed with the genetic background of Norin 29 and Shiokari, respectively. However, IR8-derived sd1 did not show effects of increasing panicle length, which was also observed with the genetic background of Koshihikari in this study. On the other hand, the results showed that Jukkoku-derived sd1 solely increased panicle length by more than 2.5%, as compared to the original cultivar Koshihikari and other semidwarfing alleles.

The first author reported transcription of the sd1 gene derived from Jukkoku [32]. RT-PCR of root RNA showed that the 779 bp fragment derived from the sd1 locus was clearly cleaved into 613 bp and 166 bp fragments by PmaCl digestion on the SNP in Jukkoku sd1, but there was no cleavage of the Sd1 locus in Koshihikari (Figure 5). This is the first evidence of the transcription of sd1, a defective gene of GA20-oxidase gene, which had reduced the culm length by the target SNP in Jukkoku sd1. Accordingly, Jukkoku sd1, which is substituted by only a single nucleotide against the 385 bp-deficient IR sd1 [17] and is actively transcribed, has no deteriorative effect on the panicle and may have a positive effect on panicle elongation by the overflow of nutrition due to culm reduction.

In Japan, Koshihikari suffers considerable lodging damage as a result of frequent powerful typhoons and, thus, the development of lodging-resistant cultivars has been a long-standing challenge. The first author transferred the semidwarf gene Jukkoku sd1 to Koshihikari to develop a semidwarf form of Koshihikari which could withstand a typhoon by backcrossing with Koshihikari eight times [33], with more than 99.8% of the background of the Koshihikari genome, except for Jukkoku sd1 [32]. This cultivar, which was about 20 cm shorter than Koshihikari, was named Hikarishinseiki (rice cultivar number 12273) [33, 34]. Hikarishinseiki is the first cultivar to be a Koshihikari-type semidwarf with sd1 registered in Japan and USA [35, 36].

In this study, a MiSeq next-generation sequencer was used to achieve 5× rice genome coverage; namely, 4.5 ng of five DNA libraries was applied in one MiSeq run. Using this approach, the targeted gene mutation in Jukkoku sd1 was successfully detected as the SNP (G to T) in the defective GA20-oxidase gene, which had reduced the culm length by a loss-of-function mutation of GA synthesis, localized at position 38,382,746 from the end of the short arm of chromosome 1 of Koshihikari. This would be the minimum scale to detect Jukkoku sd1 in practical breeding. In Japan, genetically modified organisms are not acceptable to consumers; thus the target SNP in Jukkoku sd1 would be effectively tracked by using NGS of 5× coverage scale in each backcrossed generation with Koshihikari. On the other hand, as the DNA sequence and function of sd1 have been deciphered [15, 18, 37], new breeding methods, such as RNAi gene silencing [38] and genome editing [39], are available in other countries to retard culm length by knockout of Sd1.

The effects on phenotypic traits of rice differed between the two semidwarf genes (sd1 and d60) and also between sd1 loci of different origins (Jukkoku sd1 and IR8 sd1). The effect on culm length was more pronounced in plants carrying d60 than in those carrying sd1 (culm length: Jukkoku sd1, 71.8 cm; IR8 sd1, 68.5 cm; and d60, 65.7 cm). The reduction in internode intervals was relatively uniform in lines carrying Jukkoku sd1 or d60 (uniform reduction type), while reductions of the third and lower intervals were larger than in the upper intervals in lines carrying IR8 sd1 (Figure 3 and Table 1). Thus, the center of gravity will be lower in the lower internode reduction type, suggesting higher lodging resistance in lines carrying IR8 sd1 than in those carrying Jukkoku sd1 or d60. Although the function of d60 is not known, it is clearly distinct from that of sd1, as the additive double-dwarf effect of sd1 and d60 appeared in Jukkoku sd1 plus d60 and IR8 sd1 plus d60, respectively. The results of this study demonstrated that d60 has a stronger effect than sd1 on culm length and exerts similar effects on other phenotypic traits as with sd1. Although many semidwarf genes are associated with a reduction in panicle length, d60 does not exert such negative effects on phenotypic traits of rice plants. Taken together, these results showed that d60 is useful for adding genetic diversity to semidwarf varieties and is thus of particular importance in the field of plant breeding.

Conflicts of Interest

The authors have no conflicts of interest to declare.

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Supplementary Materials

DNA sequences of Sd1/sd1 locus. (Supplementary Materials)

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