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## Research Paper

# Validation of a quantitative trait locus for the white-core expression rate of grain on chromosome 6 in a brewing rice cultivar and development of DNA markers for marker-assisted selection

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*Sake*-brewing cultivars among varieties of Japanese rice (*Oryza sativa* L.) have traits adapted to the *sake*-brewing process, such as a high white-core expression rate (WCE). Our previous study detected putative quantitative trait loci (QTLs) associated with a high WCE derived from Yamadanishiki, a popular brewing rice cultivar. Because the occurrence of white-core grains depends on air temperature and the position of the grain on the panicle, phenotyping of WCE must consider these variable conditions. In this study, *qWCE6*, a QTL for the WCE on chromosome 6, was validated for the first time, and the phenotyping method examined for its suitability in fine-mapping. A clear tendency towards high WCE was observed in late-heading substituted lines which headed under low daily mean temperature at the experimental location. White-core grains were often expressed by the primary spikelets on the upper panicle, producing a high percentage of superior grains. The segregating population for *qWCE6* in late heading revealed a distinct difference in WCE between the Koshihikari and Yamadanishiki homozygous alleles at *qWCE6* as determined from that locality. Further, two insertion/deletion markers were developed for the marker-assisted selection of *qWCE6*. Our results will be useful for informing the breeding of *sake*-brewing rice cultivars.

**Key Words:** *sake*-brewing rice cultivar, QTL, white-core grain, MAS, Yamadanishiki.

## Introduction

Japanese rice wine, “*sake*”, has recently become popular internationally, and the export amount has increased by about 2.7 times in the last 10 years (National Tax Agency Japan 2018). To produce *sake*, grains of rice (*Oryza sativa* L.) are first polished to a high grade and steamed after water absorption. Next, “*koji*” is fermented with the steamed rice and *Aspergillus oryzae*. Finally, *sake* is brewed from a combination of *koji*, a large amount of steamed rice, yeast (*Saccharomyces cerevisiae*), and water. Brewing rice cultivars are used which have been adapted to the brewing process and possess grain traits different from those of cooking rice cultivars. Of these traits, a high white-core expression rate (WCE) is characteristic of brewing rice with respect to grain appearance, and breeding of brewing rice cultivars targets this trait (Ikegami *et al.* 2003). White-core grain has an opaque structure in the central grain and displays positive

effects on water absorption and, in particular, on the fermentation of *koji* (Horigane *et al.* 2014, Yanagiuchi *et al.* 1997). Elucidation of the genetic basis of this trait is therefore of interest for brewing rice cultivar breeding.

The quantitative trait loci (QTLs) associated with WCE in Yamadanishiki, the most popular brewing rice cultivar in Japan, have been previously reported. Yoshida *et al.* (2002) detected several QTLs on chromosome 12 and at an unknown position, and Okada *et al.* (2017) detected five QTLs on chromosomes 1 (*qWCE1-1* and *qWCE1-2*), 3 (*qWCE3*), 6 (*qWCE6*), and 10 (*qWCE10*), which increased the WCE of the Yamadanishiki allele. Because the latter study used a greater number of DNA markers and lines than the former, its results should be treated with higher confidence. Of the detected QTLs, *qWCE6* was found to have the largest genetic effect, and multiple QTLs for WCE were reported around *qWCE6* in non-brewing rice cultivars (Peng *et al.* 2014, Tan *et al.* 2000). *qWCE6* can thus be regarded as a highly reliable QTL for WCE; however, the genetic effects of all the QTLs associated with WCE have not been validated to date.

Chalky grains, including white-core grains, often result from high temperatures during rice ripening (Morita 2008, Tsukaguchi *et al.* 2012, Zhao and Fitzgerald 2013), and

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some heat-induced QTLs and genes associated with chalky grain have been identified (Hakata *et al.* 2012, Kobayashi *et al.* 2007, 2013, Miyahara *et al.* 2017, Nakata *et al.* 2017, Wada *et al.* 2015). On the other hand, the WCE of brewing rice cultivars exposed to high temperatures during ripening is lower than under normal conditions (Ishii *et al.* 2008). This phenomenon is found under both natural and controlled conditions, and was determined to yield larger white-core under high temperatures (Ikegami *et al.* 2015, Ishii *et al.* 2008). Therefore, very large size white-core was judged as milky-white grain (Ishii *et al.* 2008). Increasing the incidence of grain with large white-core also leads to an unwanted increase in the number of cracked grains under high-grade polishing. In addition, Okada *et al.* (2017) also detected an interaction between QTL and annual environmental conditions for *qWCE6*, in that the genetic effect of *qWCE6* was lower in a year with a higher daily average temperature during the ripening stage. Thus, environmental conditions, in particular temperature, should be considered when evaluating WCE accuracy.

Furthermore, Nagato and Ebata (1958) speculated that there is a relationship between the occurrence of white-core grains and spikelet position on the panicle in the brewing rice cultivars Hattan-Jugo and Hyogo-Omachi. They reported that WCE was higher in the upper versus lower branches, and that the vigorously growing superior grains often expressed a white-core. The present study aimed to verify this phenomenon in Koshihikari, Yamadanishiki, and the chromosomal segment substitution lines selected for investigation.

The purpose of this study is to contribute to the breeding of brewing rice cultivars through validation of the genetic effect of *qWCE6* and development of DNA markers for marker-assisted selection (MAS) of *qWCE6*. To validate *qWCE6*, we used several backcrossed lines and residual heterozygous lines (RHLs) derived from crossing Koshihikari and Yamadanishiki, as employed by Okada *et al.* (2017). Koshihikari and Yamadanishiki respectively have early and late heading dates, with a difference of about 20 days. The heading date distinction was largely expressed by *qDTH3* (Okada *et al.* 2017, 2018a), which had very large genetic effects. Two causal genes linked closely have already been established as *Hd6* and *Hd16* (Hori *et al.* 2013, Takahashi *et al.* 2001). We were thus able to control whether plants had an early or late heading date by selecting the genotypes of DNA markers based on the causal variants of *Hd6* and *Hd16*. In the case of rice plants with an early heading date, the daily average temperature during the ripening stage was higher than for those with a late heading date under our cultivation conditions. We therefore used lines with early and late heading date because WCE of Yamadanishiki and the genetic effect of *qWCE6* were variable within an environment during the ripening stage (Ishii *et al.* 2008, Okada *et al.* 2017).

## Materials and Methods

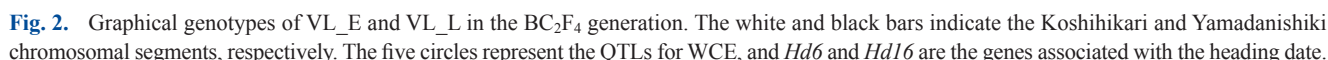
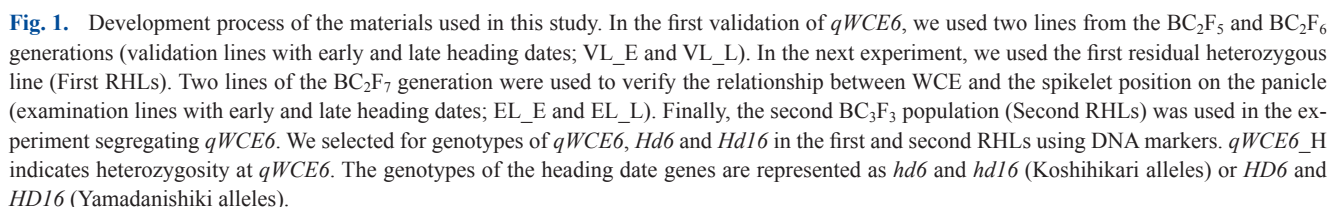
### Plant materials

**Fig. 1** shows the development process of materials used in this study. We used lines that featured a substituted chromosomal segment of Yamadanishiki on a Koshihikari genetic background. In total, two pairs of backcrossed lines and two RHLs were employed.

A total of the four backcrossed lines was derived from population increasing fixation by selfing BC<sub>2</sub>F<sub>1</sub>, generated by successively backcrossing Koshihikari twice followed by a F<sub>1</sub> crossing between Koshihikari and Yamadanishiki. These lines were genotyped with 106 DNA markers in the BC<sub>2</sub>F<sub>4</sub> generation using DNA extracted from the leaves following the approach of Dellaporta *et al.* (1983) with minor modifications (**Supplemental Table 1**). Two of the lines were used to verify the genetic effects of *qWCE6* on the BC<sub>2</sub>F<sub>5</sub> and BC<sub>2</sub>F<sub>6</sub> generation and treated as validation lines with early or late heading date (VL\_E and VL\_L; **Fig. 1**). The genetic structure of the other two lines, also selected under early or late heading date conditions, had only *qWCE6* at the Yamadanishiki allele of QTLs, as determined by our previous study, although a genotype of *qWCE3* in VL\_L remained unclear (**Fig. 2**).

The other backcrossed lines were utilized to examine the WCE evaluation method focusing on the position of spikelets on a panicle in the BC<sub>2</sub>F<sub>7</sub> generation, and to confirm whether a backcrossed line on Koshihikari genetic background also tended to express white-core on superior grains (**Fig. 1**; Nagato and Ebata 1958). These two backcrossed lines (examination line with early or late heading date; EL\_E and EL\_L) were homozygous at the *qWCE6* Yamadanishiki allele and had similar WCEs and heading date to VL\_E and VL\_L, respectively.

Two RHLs populations were used to verify the genetic effect of *qWCE6* under segregation of *qWCE6* and the heading date control, because VL\_E and VL\_L had residual chromosomal segments of Yamadanishiki on regions except for *qWCE6*. These populations were selected by MAS with RM1369 and RM2615 as the markers enclosing *qWCE6*, a cleaved amplified polymorphic sequence (CAPS) marker targeted at a functional nucleotide polymorphism (FNP) of *Hd6*, and a PCR-confronting two-pair primer (PCR-CTPP) marker targeted at an FNP of *Hd16* (**Supplemental Table 1**). One RHLs population (First RHLs) was derived from backcrossing a BC<sub>2</sub>F<sub>3</sub> generation plant with Koshihikari (**Fig. 1**). From the First BC<sub>3</sub>F<sub>2</sub> generation, we selected a rice plant for which *qWCE6* was heterozygous and *Hd16* was homozygous at the Yamadanishiki allele. *Hd6* was already fixed as homozygous at the Koshihikari allele. The other RHLs population (Second RHLs) was developed by backcrossing a BC<sub>2</sub>F<sub>5</sub> generation plant with Koshihikari and self-pollinating (**Fig. 1**). In this population, we selected a plant that was *qWCE6* heterozygous and *Hd6* and *Hd16* homozygous at the Yamadanishiki allele from the Second BC<sub>3</sub>F<sub>2</sub> generation.



VL\_E, VL\_L, and the parents were cultivated in an experimental field at Kobe University, Food Resources Education and Research Center (Kasai City, Hyogo Prefecture Japan; 34.88°N, 134.86°E) in 2013 and 2014. After harvesting, the grains were air-dried for three days and the WCE was evaluated. After dehulling, 100 grains were selected, omitting immature and cracked grains. The grains were visually inspected with a light box, white-core grains were counted, the WCE was calculated, and the occurrence rates

The greater part of the difference in heading date between Koshihikari (Aug. 3 in both years) and Yamadanishiki (Aug. 20 in 2013 and Aug. 17 in 2014) was driven by the tight linkage between *Hd6* and *Hd16* alleles (Fig. 2; Okada *et al.* 2017, 2018a).

### Confirmation of *qWCE6* genetic effects using RHLs

The First RHLs were cultivated in 72 plants. We sampled leaves from all plants and extracted DNA. To derive the *qWCE6* genotype with the two markers (RM1369 and RM2615), we measured the WCE of nine (Koshihikari homozygous), 21 (heterozygous) and six (Yamadanishiki homozygous) plants, respectively. The WCE of the First RHLs was calculated by measuring the number of white-core grains in 100 grains using the same method as described above.

The Second RHLs were cultivated in 96 plants, and the *qWCE6* genotype was identified with the same markers as above. We selected 19 (Koshihikari homozygous), 23 (heterozygous) and 15 (Yamadanishiki homozygous) plants, respectively. The WCE evaluation method was determined by the results of the experiment described below (see next section). We thus measured each WCE using 20 grains sampled from the primary spikelets (PS) on the upper part of one panicle per plant (Fig. 3). During sampling, superior grains were selected preferentially. Furthermore, dominance degree value (DDV) of *qWCE6* was calculated according to Ukai (2000), namely the absolute value of dominance effect divided by additive effect of the QTL. In addition, in five samples selected randomly from each genotype population of *qWCE6* we measured grain length and width using SmartGrain 1.0 (Tanabata *et al.* 2012).

### Examination of the WCE evaluation method focusing on the position of spikelets on a panicle

Heading dates of EL\_E and EL\_L were also similar to

VL\_E and VL\_L, respectively (EL\_E: Aug. 3, 2015 and EL\_L: Aug. 26, 2015). For five panicles of each line and parent, the position of the spikelets was registered, and we checked whether each spikelet had a white-core. Next, the WCE was calculated for each part of the panicle, including PS, secondary spikelet (SS), upper-panicle PS and lower-panicle PS (Fig. 3).

### Statistical analysis

We used R for the statistical analyses (ver. 3.4.1, R Core Team 2017). A Dunnett's multi-comparison test was used for first validation. A Steel-Dwass test was conducted for experiments except for the first validation, using the NSM3 package (Schneider *et al.* 2018). Differences in grain length and width of Second RHLs were assessed using a Tukey-Kramer test.

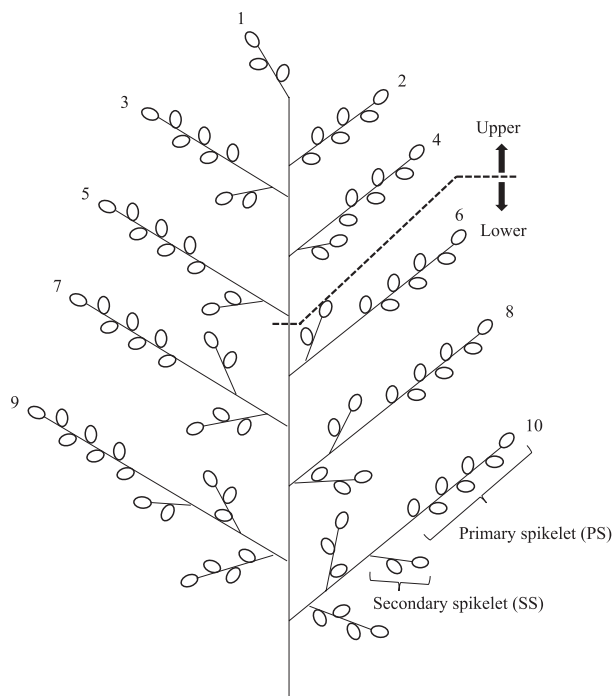
### Development of insertion/deletion (INDEL) markers

First, we searched INDEL sites using rice TASUKE (Kumagai *et al.* 2013, <https://ricegenomes.dna.affrc.go.jp/>), restricting the search to sequence depth values >5 between 1,534,301 bp and 3,662,323 bp on chromosome 6 on Koshihikari (ID: SAMD00045947) and Yamadanishiki (ID: SAMD00010633), using IRGSP-1.0 (Kawahara *et al.* 2013). The candidate region of *qWCE6* was selected based on a previous QTL analysis of WCE (Okada *et al.* 2017, 2018b). We developed two INDEL markers, verified under the following PCR conditions: 1 cycle at 95°C for 2 min, 35 cycle at 95°C for 30 s, at 57°C for 30 s and 68°C for 1 min, post-extension at 72°C for 5 min, and stop at 4°C; using a reaction medium of 2 µl of crude extracted DNA, 3 µl of 2 µM mixed primer and 5 µl of Quick Taq™ HS DyeMix (Toyobo Life Science). Amplified products were separated by electrophoresis on a 3 % agarose gel and stained with ethidium bromide.

## Results

### Validation of the genetic effect of *qWCE6*

The graphical genotypes of VL\_E and VL\_L were shown in Fig. 2. VL\_E had Koshihikari homozygous on both *Hd6* and *Hd16* and headed on Jul. 28, 2013 and Jul. 26, 2014, while VL\_L had Yamadanishiki homozygous at both *Hd6* and *Hd16* and headed on Aug. 21, 2013 and Aug. 20, 2014 (Table 1, Fig. 2). The WCE of VL\_E did not significantly differ from that of Koshihikari, while the WCE of VL\_L was significantly higher in 2013 (Table 1). In contrast, in 2014 the WCEs of both lines were significantly higher than that of Koshihikari (Table 1). This confirmed the genetic effect of *qWCE6* but left a need for additional experiments, as both lines still contained large chromosomal segments of Yamadanishiki. In addition, the WCE of VL\_E was variable in both years, while the WCE of VL\_L was comparatively stable (Table 1).



**Fig. 3.** Sampling panicle position. We classified positions on the panicle into “upper” and “lower” as follows: half the number of total primary branches counted from the top of the panicle were designated as “upper”, and the other half was designated as “lower”.



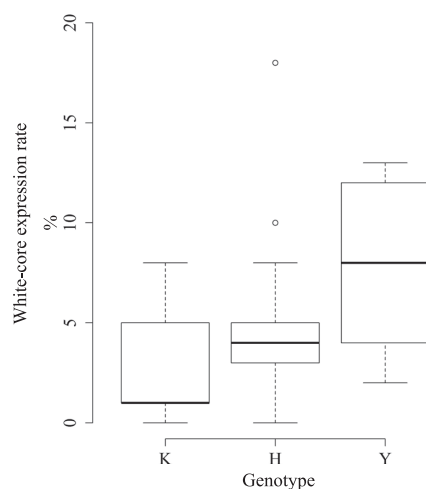
**Table 1.** Validation of the genetic effects of *qWCE6*

	WCE (%) <sup>a</sup>		Days to heading <sup>b</sup>	
	2013	2014	2013	2014
VL_E	9.4 ± 9.1	29.0 ± 6.8**	94 ± 1	95 ± 1
VL_L	18.7 ± 5.3**	34.3 ± 1.9**	118 ± 0	120 ± 1
Koshihikari	5.0 ± 2.8	9.7 ± 2.7	100 ± 1	102 ± 1
Yamadanishiki <sup>c</sup>	70.5 ± 4.2	70.3 ± 4.1	117 ± 0	116 ± 1

<sup>a</sup> Phenotypic data indicate “mean ± standard deviation (SD)”, and the asterisks represented significant difference between Koshihikari and each line ( $P < 0.01$ ).

<sup>b</sup> Significant difference test was not conducted for the data.

<sup>c</sup> Yamadanishiki's data are supplemental values.



**Fig. 4.** WCE classified the First RHLs into *qWCE6* genotypes. On the horizontal axis, K, H, and Y represent the *qWCE6* genotype for Koshihikari homozygous, Koshihikari heterozygous, and Yamadanishiki homozygous, respectively.

### First segregation test of *qWCE6*

The heading date of the First RHLs was from Aug. 14–24, and the population included late or slightly late heading lines. The heading date data for each genotype population of *qWCE6* were Aug. 19–22 (Koshihikari homozygous), Aug. 15–23 (heterozygous) and Aug. 14–16 (Yamadanishiki homozygous). This distribution of heading dates resulted from segregating *Hd17* in the candidate region of *qWCE6* (Matsubara *et al.* 2012, Okada *et al.* 2017, 2018a). WCE of First RHLs was evaluated using the conventional method, but no statistically significant difference was detected among *qWCE6* genotypes (Fig. 4). The lines that were homozygous at the *qWCE6* Yamadanishiki allele appeared to have a higher WCE, but this tendency was not found to be statistically significant (Fig. 4). It was therefore necessary to consider improving the method used to evaluate WCE for *qWCE6* validation and fine-mapping. As white-core grains generally express at superior grains on a panicle (Nagato and Ebata 1958), this was done by focusing on the spikelet position on a panicle, as described in the next section.

**Table 2.** Relationship between WCE and spikelet position on the panicle

	WCE of partial panicle (%) <sup>†</sup>			
	PS <sup>‡</sup>	SS <sup>‡</sup>	Upper PS	Lower PS
EL_E	30.2 ± 13.2 <sup>a</sup>	7.7 ± 3.4 <sup>a</sup>	33.5 ± 12.3 <sup>ab</sup>	26.2 ± 19.1 <sup>a</sup>
EL_L	28.7 ± 5.4 <sup>a</sup>	8.1 ± 5.3 <sup>a</sup>	45.0 ± 9 <sup>b</sup>	10.4 ± 10.1 <sup>a</sup>
Koshihikari	19.9 ± 7.6 <sup>a</sup>	5.6 ± 2.7 <sup>a</sup>	20.4 ± 9.3 <sup>a</sup>	19.5 ± 9.7 <sup>a</sup>
Yamadanishiki	85.2 ± 5 <sup>b</sup>	65.7 ± 7.2 <sup>b</sup>	85.4 ± 7.3 <sup>c</sup>	85.1 ± 8 <sup>b</sup>

<sup>†</sup> WCE data show mean ± SD; the different letters in supertext represent statistical difference between groups (Steel-Dwass method,  $P < 0.05$ ).

<sup>‡</sup> PS and SS represent primary and secondary spikelet, respectively.

### Relationship between WCE and spikelet position on a panicle

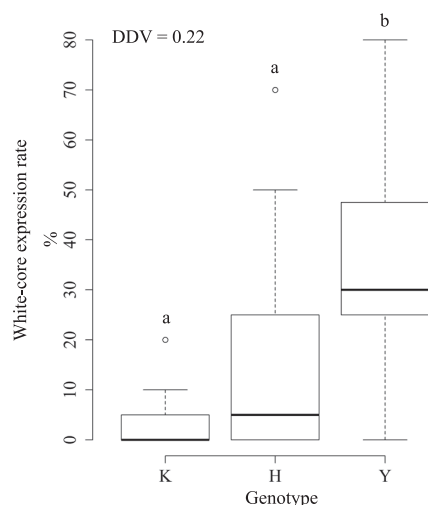
We first inspected the WCE of PS and SS (Table 2). PS WCE was higher than that of SS for all materials, but the WCEs of spikelets in both EL\_E and EL\_L were not higher than in Koshihikari (Table 2). We next focused on the PS, which had a high WCE, and divided the measure into PS on the upper panicle and lower panicle, respectively. EL\_L was found to have a significantly increased PS WCE on the upper panicle (Table 2). This finding suggests that it is important to evaluate PS WCE on the upper panicle in late-heading material, sampling as many superior grains as possible.

### Second segregation test of *qWCE6*

The heading dates of the Second RHLs were from Aug. 23 to Sep. 1, and late-heading was a characteristic of the heading date for this population. The heading date data for each genotype population of *qWCE6* was from Aug. 25 to Sep. 1 (Koshihikari homozygous), Aug. 23–27 (heterozygous) and Aug. 23–27 (Yamadanishiki homozygous). The segregation of Second RHLs also was due to *Hd17* (Matsubara *et al.* 2012, Okada *et al.* 2017, 2018a). We evaluated WCE using 20 grains from the PS on the upper panicle (superior grains as priority) based on the results of the above experiment. We found that the WCE of the population that was homozygous at the Yamadanishiki allele of *qWCE6* was significantly higher than that of the Koshihikari-homozygous population (Fig. 5). This indicates a better performance of this evaluating method than that of the method used for the First RHLs, although rice plants with a low WCE also existed in the Yamadanishiki-homozygous population. At the same time, the DDV of *qWCE6* was 0.22 (Fig. 5). Grain length and width did not change with *qWCE6* genotype (Supplemental Fig. 1).

### Development of INDEL markers

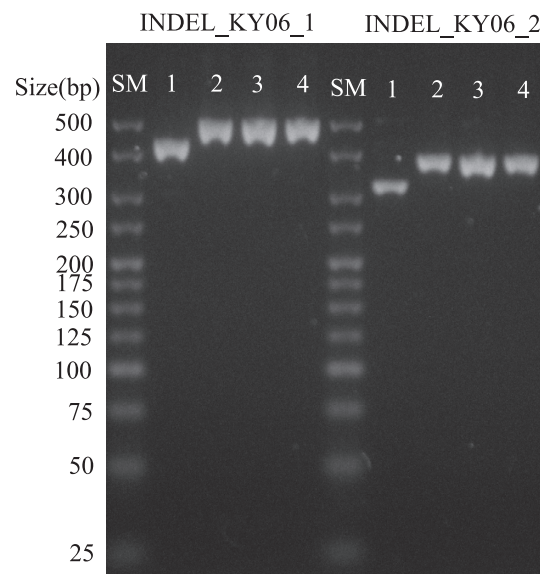
A total of 14 INDEL sites was searched, to which one site at 3,652,857 bp was added due to its close proximity to one upstream INDEL site (Supplemental Table 2). Among these 15 sites, we targeted INDELs of 2,376,346 bp and both 3,652,851 bp and 3,652,857 bp because of relatively large INDELs. Two INDEL markers (INDEL\_KY06\_1 and INDEL\_KY06\_2) were established (Table 3, Fig. 6).



**Fig. 5.** WCE classified the Second RHLs into *qWCE6* genotypes. On the horizontal axis, K, H, and Y represent the *qWCE6* genotype for Koshihikari homozygous, Koshihikari heterozygous, and Yamadanishiki homozygous, respectively. The different letters above each box-plot indicate the statistical differences between groups (Steel-Dwass test,  $P < 0.05$ ). DDV represents dominance degree value calculated as an absolute value of dominance effect divided by additive effect of a QTL. When the value is  $DDV = 0$ ,  $0 < DDV < 1$ ,  $DDV = 1$  or  $1 < DDV$ , inheritance manner of a QTL is defined as no dominance, incomplete dominance, complete dominance or overdominance, respectively (Ukai 2000).

## Discussion

In this study, a line with the QTL for WCE showed WCE variation across two years; the early heading line (VL\_E) exhibited variation in both years (2013 and 2014; **Table 1**), similar to the trend observed by Okada *et al.* (2017). In 2013 and 2014, the daily average temperature in the middle of August 2013 was higher than that in 2014 (**Supplemental Fig. 2**, cited from Okada *et al.* 2017). On the other hand, the average daily temperature from the end of August tended to show lower and lower transition in both years. This suggests that WCE variation in VL\_E resulted from differences in average daily temperatures during the middle of August, which corresponds to the ripening stage of the VL\_E. On the other hand, it is likely that the WCE of VL\_L was stable because the ripening stages of VL\_L in 2013 and 2014 were at lower temperature and under better conditions for the expression of white-core.



**Fig. 6.** Electrophoregram of insertion/deletion (INDEL) markers developed in this study. The lanes of SM, 1, 2, 3 and 4 represented size marker, Koshihikari, Yamadanishiki, VL\_E and VL\_L.

High temperatures during ripening have been shown to cause larger white-core size in a brewing rice cultivar (Ikegami *et al.* 2015, Ishii *et al.* 2008). Under the same conditions, the incidence of milky-white and white-back grains also increased in cooking rice cultivars (Ikegami *et al.* 2015). In the present study, VL\_E in 2013 showed an increased occurrence rate over VL\_L of other types of chalky grain through heat-stress, because its ripening period was at high daily mean air temperature (**Supplemental Table 3**, **Supplemental Figs. 2, 3**). We also suggest that distinguishing between white-core and chalky grains induced by heat-stress is difficult because part of the white-core fuses with other grain components in chalky grains.

We investigated the relationship between spikelet position on the panicle and WCE. As we found that there generally were many superior grains in the PS of the upper panicle, our results are consistent with those of Nagato and Ebata (1958), and we confirmed the same phenomenon for backcrossed lines on a Koshihikari genetic background. However, white-core grains found in Koshihikari and EL\_E may result from high temperatures during the ripening stage due to the difficulties in distinguishing them from other chalky grain types, as noted above. Moreover, Yamadanishiki

**Table 3.** Primers for insertion/deletion markers for marker-assisted selection

Marker name	Target position on chr.6	Forward	Reverse	Amplicon size (bp) <sup>a</sup>		AT (°C) <sup>b</sup>
				Koshihikari	Yamadanishiki	
INDEL_KY06_1	2,376,346	AAGCCAGCAGTTTGTACACATC	AAACATTAGCACACGACGAGTC	430	502	57
INDEL_KY06_2	3,652,851 and 3,652,857	ATCCCGAGTTCTCTCCGAAAC	CGCCAGAAAGAGAGAAGAGAAG	321	382	57

<sup>a</sup> Each PCR product size expected from rice TASUKE (Kumagai *et al.* 2013, <https://ricegenomes.dna.affrc.go.jp/>).

<sup>b</sup> Annealing temperature.

also had a higher WCE than the other lines in the SS (Table 2). It may be considered that high WCE in the SS resulted from having plural factors for occurring white-core because reported QTLs for WCE had low phenotypic variance explained and frequency of a line having WCE as high as Yamadanishiki was very low (Okada *et al.* 2017). Therefore, it is possible for a line with a high WCE in the SS to be developed, although the correlation between WCE in the SS and WCE evaluated with the conventional method needs to be confirmed using a segregating population.

The WCE of the First RHLs displayed no significant differences among the *qWCE6* genotypes, while the WCE of the Second RHLs was clearly distinguished between populations that were homozygous at the Koshihikari and Yamadanishiki alleles by *qWCE6* (Figs. 4, 5). In the experiment using the First RHLs, we measured WCE in 100 normal grains (randomly selected ignoring panicle position), and the genetic effect of *qWCE6* was unclear. In addition, the WCE of the population that was *qWCE6* homozygous at the Yamadanishiki allele was lower than that of VL\_E and VL\_L (Table 1, Fig. 4). To account for this phenomenon, we propose that the rate of Yamadanishiki chromosomal segments in the First RHLs was lower than in VL\_E and VL\_L due to triple backcrossing. Heterozygous regions in the population would thus still exist, as the genes increasing WCE are considered to be of slightly recessive inheritance (Kamijima *et al.* 1981). Okada *et al.* (2017) reported four QTLs that exclude *qWCE6*; however the genetic variation of the WCE population used in that study was not fully expressed by these QTLs, suggesting that other QTLs exist and have very small genetic effects.

The genetic effect of *qWCE6* was thus clearly validated under the improving sampling method employed with the Second RHLs, and white-core grain associated with *qWCE6* was found to be independent of grain size (Fig. 5, Supplemental Fig. 1). From the DDV of *qWCE6*, the mode of inheritance appears to be incomplete dominance (Fig. 5). However, some plants were found to have a low WCE despite being homozygous at the *qWCE6* Yamadanishiki allele, probably because we calculated the WCE using only 20 grains in the Second RHLs. WCE should therefore be evaluated using a greater number of grains to attain a more accurate measurement.

The developed INDEL markers were able to distinguish between Koshihikari and Yamadanishiki (Fig. 6). These INDELs at 2,376,346 bp and 3,652,851 bp on chromosome 6 showed the different alleles from Yamadanishiki in more than 80% of 215 accessions of all *temperate japonica* in the rice TASUKE. This indicates that these markers will be of ready use for MAS of *qWCE6*.

Many QTLs associated with chalky rice grain have been reported around *qWCE6*. These were detected under either ambient or high temperature conditions. The QTLs for white-back and basal-white grains induced by heat-stress were identified by Kobayashi *et al.* (2007) and Kobayashi *et al.* (2013). In contrast, Tan *et al.* (2000) and Peng *et al.*

(2014) reported QTLs for white-core and milky-white grain around *qWCE6* under ambient temperature conditions. This indicates that *qWCE6* may correspond to the latter QTLs due to the similar temperature environment. Zhao *et al.* (2016), Gong *et al.* (2017), and Qiu *et al.* (2017) detected the QTLs for the chalky grain rate around *qWCE6* and suggested that *qWCE6* is also related to these QTLs, although they did not classify chalky grain types. It has been shown that the mutation of genes associated with starch synthase induces chalky grains (Fujita *et al.* 2007, Nishi *et al.* 2001). For example, a mutation of *Isoamylase 1* on chromosome 8 was shown to increase the chalky grain rate (Sun *et al.* 2015). Some genes associated with starch synthase are present in the endosperm around *qWCE6*, such as *Waxy* and *starch synthase I (SSI)* (Yamakawa *et al.* 2008). Rice TASUKE demonstrated that two variants of *Waxy*, one each for Koshihikari and Yamadanishiki, existed in promoter region and 3' UTR as one SNP and one INDEL, respectively. On the other hand, *SSI* is invariant between Koshihikari and Yamadanishiki. The *Waxy* variants are unlikely to represent causal variants of *qWCE6* because Nipponbare (a cooking rice used as the reference sequence) and Yamadanishiki had the same allele. Further studies employing methodologies such as fine-mapping are therefore needed.

In conclusion, the present study for the first time validated the genetic effects of a white-core grain QTL, *qWCE6*, which constitutes a very important factor for Yamadanishiki. The INDEL markers developed in this study allow for MAS of *qWCE6*. We also improved WCE evaluation based on temperature and spikelet position on the panicle for use in future fine-mapping. Our findings suggest that late-heading lines should be used for evaluation at our study location, and that WCE should be measured using on the order of 100 superior grains. However, late-heading lines may only be suitable for locations comparable to that used in this study. Daily mean air temperature during the ripening stage in our location was normally below 26°C (Supplemental Fig. 2), which likely prevents white-cores from attaining great size and makes the expression of other types of chalky grains difficult (Ikegami *et al.* 2015, Ishii *et al.* 2008). In other regions, adjustments should be made to the heading date based on the environment (such as temperature) and the grain quality of the brewing rice cultivar in question. The heading date decision is thus also a key factor for breeding brewing rice cultivars. In future studies, an element of the genetic basis of white-core development will be elucidated by map-based cloning of *qWCE6*. Moreover, QTLs associated with white-core grain were also detected using mapping populations crossing between cooking rice cultivars around *qWCE6* (Peng *et al.* 2014, Tan *et al.* 2000). Therefore, an allele decreasing WCE of *qWCE6* may be available at breeding of a cooking rice cultivar having high-quality grain.



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