RESEARCH ARTICLE

PLANT SCIENCE

# Enhancing rice panicle branching and grain yield through tissue-specific brassinosteroid inhibition

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C d d b d - b
d b . B d (BR ) b
. B d d- BR
b BRASSINOSTEROID-DEFICIENT DWARF3 (BRD3) d b .
b BR b G K3/ HAGG -LIKE KINA E2
d b O MAD d TERMINAL FLOWER1
RICE CENTRORADIALIS2. BRD3 d b
b d d. O
d d
d

Rice (*Oryza sativa*) is an essential staple crop, feeding more than half of the world's population (*1*). With global population growth and diminishing arable land, it is crucial to enhance rice grain yield per unit area to ensure global food security. During the development of the rice panicle, the inflorescence meristem produces multiple primary and secondary branches. Panicle branching determines grain number, one of the determinants of grain yield. Therefore, one promising approach to increase grain yield is by enhancing panicle branching. The identification of genes controlling panicle branching, such as *Grain number1a* (*Gn1a*) and *DENSE PANICLE 1* (*DEP1*), has exemplified the success of this approach (*2*, *3*). However, after extensive breeding efforts using these genes, rice yield has reached a plateau, primarily attributed to the intricate trade-offs among various traits that prove resistant to conventional breeding interventions. Tackling this formidable challenge requires an exploration of pioneering genetic reservoirs sourced from a wide array of germplasm resources to facilitate subsequent advancements.

Brassinosteroids (BRs) are a class of steroid phytohormones that play diverse roles in plant growth and development (*4*–*6*). BR responses require BR signaling, in which GSK3/SHAGGY-like kinases, such as BRASSINOSTEROID INSENSITIV2 in the model plant *Arabidopsis* (*Arabidopsis thaliana*) and GSK3/SHAGGY-LIKE KINASE2 (GSK2) in rice, act as the central inhibitors (*7*, *8*) targeting different substrates to modulate various BR responses (*5*, *9*). In cereal crops such as rice, BRs regulate important agronomic traits such as plant height, leaf angle, and grain size (*5*). Rice mutants with defective BR synthesis or signaling usually show dwarfism, upright leaves, and small grains (*10*–*12*). Despite the promising results achieved in major crops such as rice, maize (*Zea mays*), and wheat (*Triticum aestivum*) using BR-related gene resources for yield improvement (*13*–*15*), the small-grain phenotypes limit the utility of BR mutants for enhancing grain yield. In addition, it remains largely unclear how BRs regulate panicle branching and grain number. Together with additional potential influences such as stress tolerance, the pleiotropic effects of BRs present great challenges for the effective utilization of BR regulatory genes (*5*).

Unlike the common single-grained rice, clustered-spikelet rice (CL), also known as compound-spikelet rice, wheat-spike rice, or SAN-LI-QI (meaning triple-grain-miracle in Chinese), is an unusual rice germplasm possessing an extraordinary trait of multiple (often triple) complete spikelets or grains clustered on the panicles. Since its description early in 1931, the distinctive character of CL has been used to construct a genetic linkage map by worldwide rice geneticists (*16*–*19*). Owing to its potential for improving grain productivity, researchers have made numerous efforts to identify and clone the gene (*CL*) responsible for this trait. Although several studies have successfully mapped the *CL* locus to a specific region on chromosome 6 (*20*–*24*), the gene itself has eluded cloning attempts, for reasons that are unclear.

Here, we report our successful cloning of the causal gene for CL by large-scale screening of CL suppressor mutants. We show that CL is associated with complex chromosome structural variations, which activate the expression of the BR catabolic gene *BRASSINOSTEROID-DEFICIENT DWARF3* (*BRD3*) in the secondary branch meristems and pedicels. Notably, the spatial-specific activation of *BRD3* promotes grain number while avoiding the commonly seen negative effects of BR deficiency on grain size and grain quality. Moreover, we establish a complete BR pathway for mediating BR regulation of panicle branching. Furthermore, we demonstrate that introduction of *CL* into various backgrounds, either containing *Gn1a* gene or not, can substantially improve rice yield.

## Results

### CL confers enhanced panicle branching and grain yield

CL1 represents a typical CL variety characterized by clustered growth, primarily consisting of three grains. To facilitate comparisons, we generated a non-CL variety (NCL1) by using CL1 as the recurrent parent. Apart from the distinct panicle morphology, no other notable differences were observed between NCL1 and CL1 (Fig. 1, A and B). The clustered growth in CL1 predominantly occurred at the terminus of the secondary branch, where a pair of closely fused spikelets formed, with a third spikelet attached nearby (Fig. 1, C and D). Additionally, the tertiary branch was frequently observed in CL1 but not in NCL1 plants (Fig. 1E). Moreover, pedicels attached to the spikelets in CL1 were shorter than those of NCL1 (Fig. 1F).

CL1 showed a 28.2% increase in grain number per panicle compared to NCL1 (Fig. 1G), which was attributed to a greater number of secondary panicle branches (35.2% increase) and associated spikelets, but not the primary branches (Fig. 1, H to J). No significant differences were detected between NCL1 and CL1 for other traits such as panicle length, panicle number, heading date, and 1000-grain weight (Fig. 1, K to M, and fig. S1). The specific enhancement of grain number in CL1 resulted in a 20.1% increase in grain yield per plant in Beijing (north of China, 2021), 21.7% increase in Sanya (south of China, 2022), and 12.8% increase in Fuzhou (south of China, 2020) compared to NCL1 (Fig. 1, N and O, and fig. S1). These findings were further confirmed in the field plot tests. Under the field conditions, except for the CL phenotype, no other differences were observed throughout NCL1 and CL1 growth (Fig. 1P). Compared to NCL1, CL1 grain yield per hectare averaged 20.96, 17.99, and 11.27% higher in Beijing, Sanya, and Fuzhou, respectively (Fig. 1Q and fig. S1). Regarding grain quality, no significant differences were detected between NCL1 and CL1 in terms of grain chalkiness, amylose content, gel consistency,

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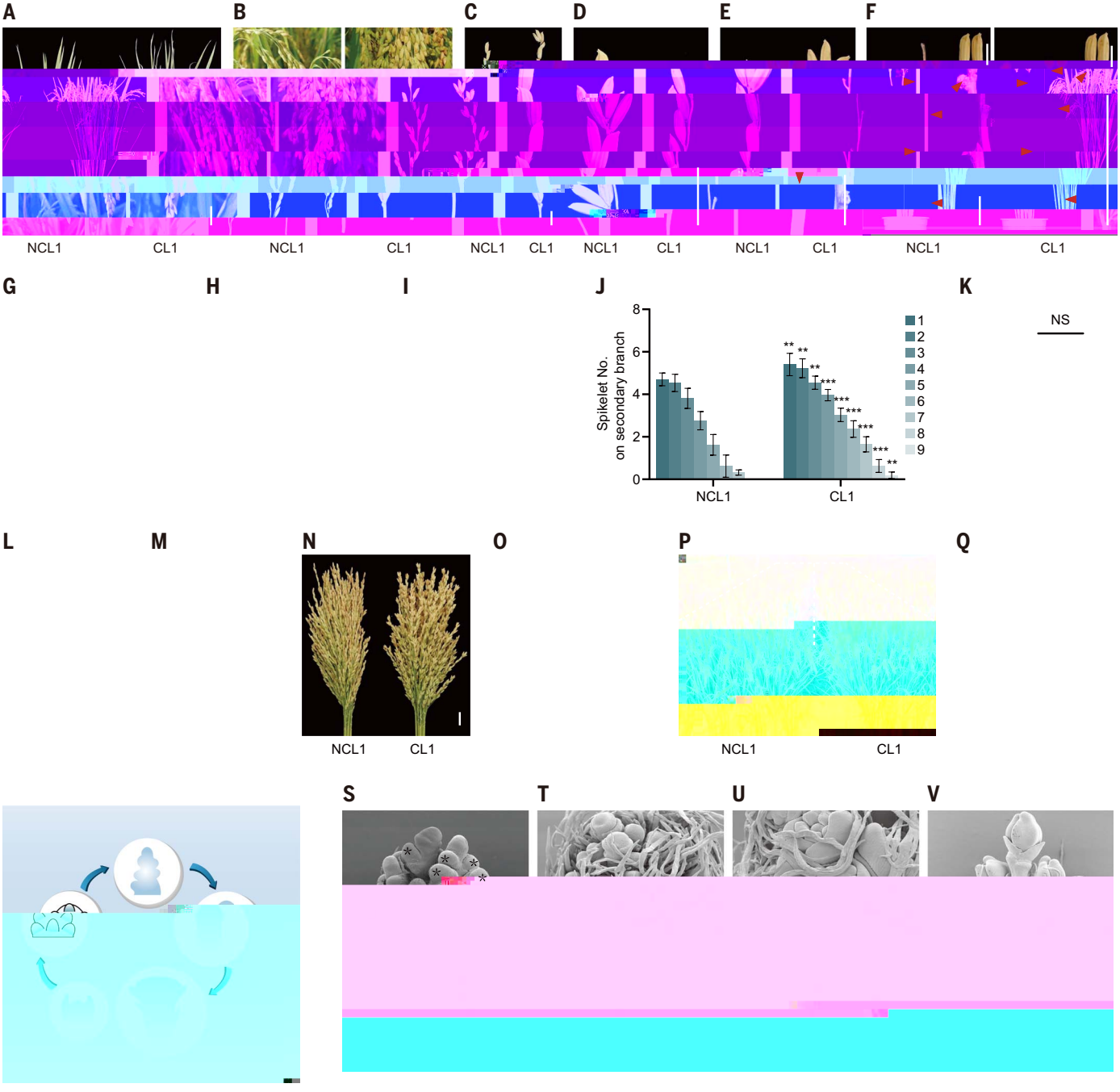
†These authors contributed equally to this work.




and gelatinization temperature, and only total protein content was slightly greater in CL1 (fig. S2). These results suggested that the CL trait has great potential for crop improvement, prompting us to explore the underlying mechanisms.

## Prolonged branching time leads to increased grain number

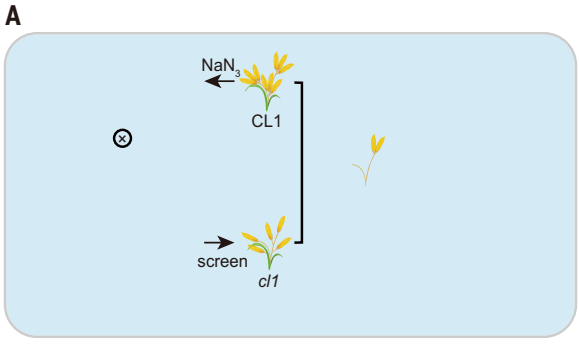
During rice panicle development, a series of lateral meristems with distinct identities are sequentially generated (Fig. 1R). Inflorescence meristems first produce lateral primary branch meristems (PBMs). Subsequently, the newly initiated meristem from the PBM either acquires the identity of secondary branch meristem (SBM) or terminates as a spikelet meristem (SM). The SBM continues to generate next-order lateral meristems that will eventually

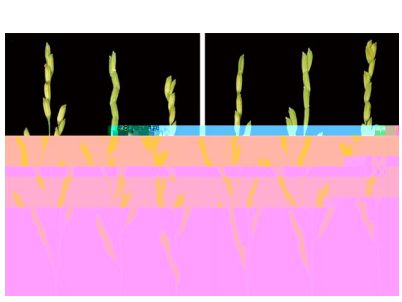
acquire the SM identity. Finally, the SM is transformed into a floral meristem (FM) (*25*–*27*). Scanning electron microscopic (SEM) observation identified excrescent SBMs in CL1 at the early stage of panicle development (fig. 1S). In addition, the transition from BM to SM was delayed in CL1 (Fig. 1, S to U). FMs were formed in NCL1 when the young panicle of CL1 was at the SM stage (Fig. 1T), and the FMs were already fully developed in NCL1 when the FMs in the young panicle of CL1 were formed (Fig. 1U). The delay in the transition from BM to SM resulted in prolonged time for generating more SBMs and even the formation of tertiary branches in CL1 (Fig. 1E). Furthermore, an additional SM developed adjacent to the terminal spikelet (Fig. 1U). Pedicel elongation was clear in NCL1 but barely observed in CL1 (Fig. 1V). On the basis of these observations, we concluded that three developmental phenomena drive CL formation: (i) the development of more SBMs, (ii) the initiation of supernumerary SM, and (iii) shortened pedicels.

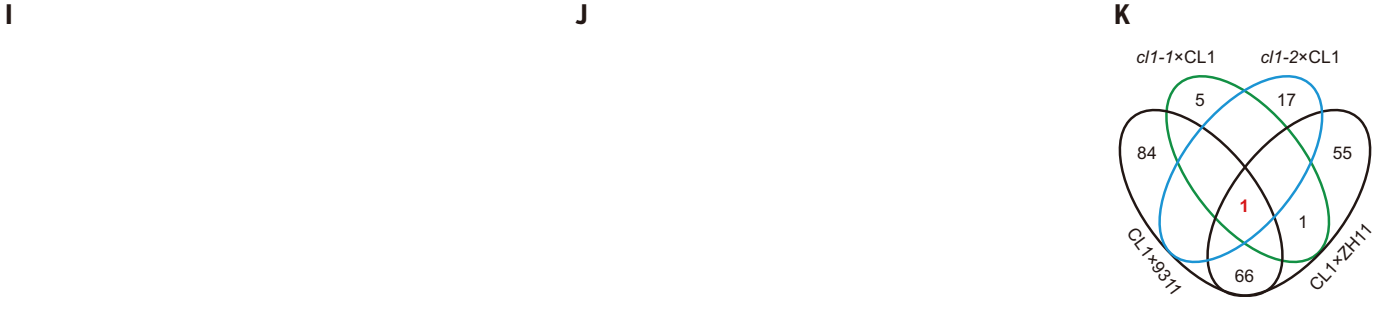
### Cloning the causal gene through screening CL suppressors

To help us track down the genetic basis for CL, we requested three other CL germplasms (CL2, CL3, CL4) from different sources, including one (CL3, IRIS_313-11403) from the 3000 Rice Genomes Project (3K-RG) (*28*). All three showed the typical CL phenotype, but they exhibited highly variable plant architecture and panicle morphology (fig. S3). We crossed them with each other and found that all $F_1$ plants derived from the different crosses retained the CL phenotype (fig. S3). By contrast, $F_1$ plants derived from crosses between CL1 and 9311, an NCL *indica* variety, or between CL1 and Zhonghua11 (ZH11), an NCL *japonica* variety, showed only a weak CL phenotype (WCL), characterized by clustered growth of two grains at the terminus of the panicle branches (figs. S4 and S5). In addition, in the crossed CL1 × ZH11 $F_2$ population, the segregation ratios (NCL:WCL:CL) were 65:131:45 (fig. S5), close to 1:2:1 ($\chi^2$ = 0.0951664 < 5.99 at $P$ = 0.05). These results suggested that the distinctive phenotype of these CL plants is determined by the same semidominant locus. However, our intense efforts using either map-based cloning or bulked-segregant analysis (BSA) by generating large populations or different crosses could only locate the locus to the ~22.85- to 23.85-Mb (megabase) region on chromosome 6 (fig. S6), like previously reported attempts (*21*–*24*). One possibility is that there exist complex chromosome structural variations around the region.

The failure of these routine cloning methods prompted us to design a strategic cloning roadmap by generating large-scale mutagenesis of CL1 for identifying the causal gene (Fig. 2A). From a population of 10,000 independent mutant lines ($M_1$) using sodium azide as the mutagen, examination of the phenotypes of the $M_2$ plants (16 for each line) led to the identification of two mutant lines that reverted to a wild-type NCL phenotype, which we designated *cl1-1* and *cl1-2* (Fig. 2B and fig. S7). By comparing the NCL and CL plants pooled from the $F_2$ population of the *cl1* × CL1 backcrosses, we found that, similar to NCL1, both *cl1-1* and *cl1-2* had significantly decreased grain number compared to CL1, owing to their smaller numbers of secondary branches (Fig. 2, D to F). Both mutants also showed identical plant architecture, panicle number, panicle length, and 1000-grain weight to those of NCL1 and CL1 (figs. S8 and S9 and Fig. 2G). In addition, the grain yield per plant of the two *cl1* mutants decreased by 16.4% to 27.7% compared to CL1 in Sanya (Fig. 2H). Because CL is controlled by a single semidominant



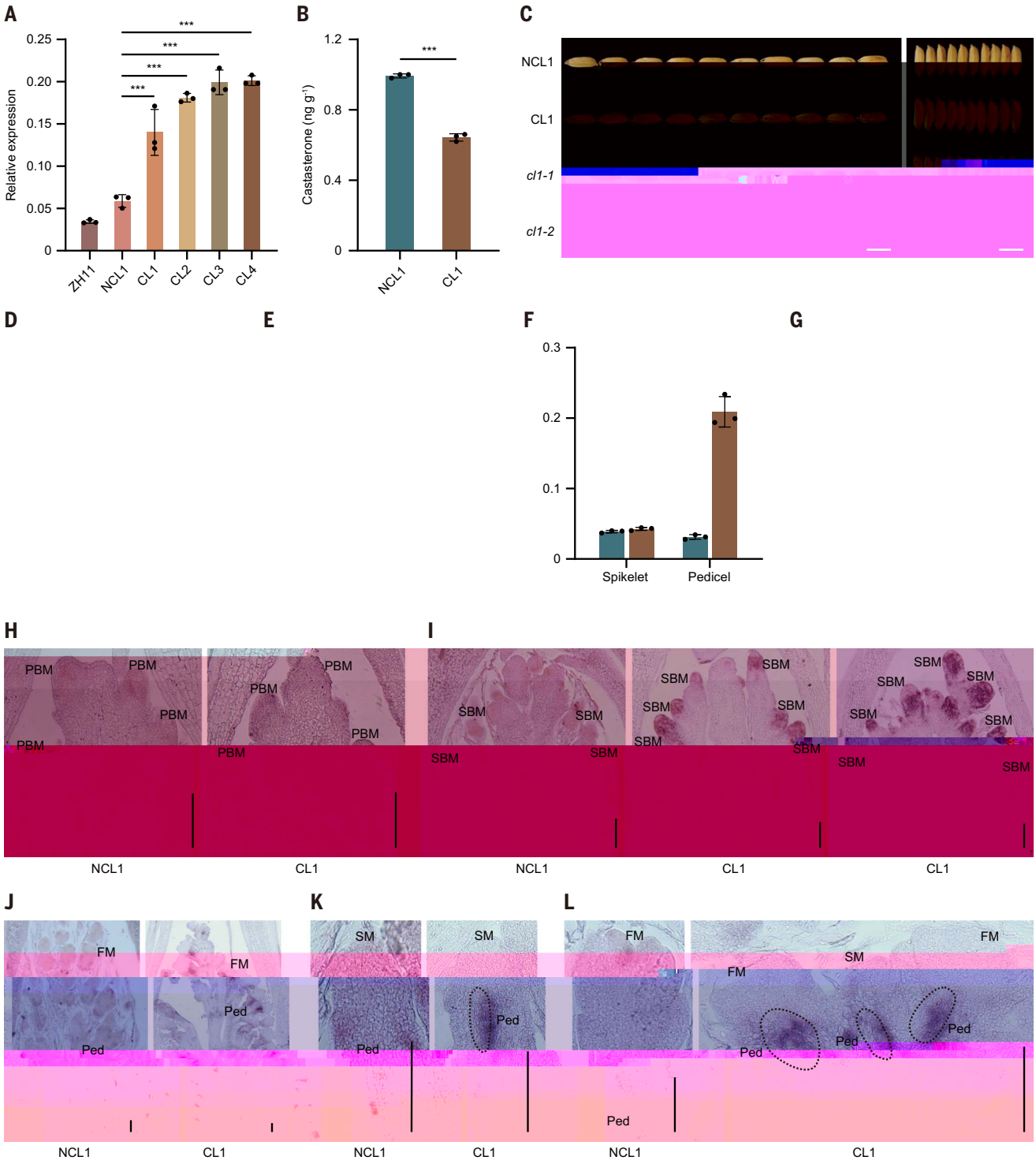





no detectable differences in grain size and plant height among CL1, *cl1-1*, *cl1-2*, and NCL1 (Fig. 3, C to E, and fig. S17). Because CL is closely associated with shortened pedicels, we first compared the expression pattern of *BRD3* in pedicels and spikelets of NCL1 and CL1. Notably, *BRD3* was up-regulated in pedicels, but not spikelets, of CL1 (Fig. 3F). To confirm this result, we also measured BR content in these tissues of both lines. Consistent with the expression pattern of *BRD3*, the castasterone level was only ~20% lower in CL1 spikelets, yet ~70% lower in CL1 pedicels, compared to NCL1 spikelets and pedicels (Fig. 3G).

We further conducted RNA in situ hybridization to determine the spatiotemporal expression pattern of *BRD3* during panicle development. During the development of NCL1 inflorescences, *BRD3* expression was very weak and not specific to any tissues (Fig. 3, H and I). By contrast, we detected specific signal with


A
Relative expression
0.25
0.20
0.15
0.10
0.05
0
***
***
***
***
ZH11
NCL1
CL1
CL2
CL3
CL4
B
Castasterone (ng g-1)
1.2
0.8
0.4
0
***
NCL1
CL1
C
NCL1
CL1
cl1-1
cl1-2
D
E
F
0.3
0.2
0.1
0
Spikelet
Pedicel
G
H
PBM
NCL1
CL1
I
SBM
NCL1
CL1
CL1
J
FM
Ped
NCL1
CL1
K
SM
Ped
NCL1
CL1
L
FM
SM
Ped
NCL1
CL1

top-down development in SBM, but not PBM, of CL1 panicles (Fig. 3, H and I). At the stage of SM or FM development, *BRD3* expression was specifically detected at the base of spikelets, corresponding to the pedicel positions, in CL1 but not NCL1 (Fig. 3, J to L). As a negative control, no signal was detected when *BRD3* was used as the sense probe (fig. S18). The strong relationship between *BRD3* expression pattern and plant phenotype suggests that spatial enhancement of *BRD3* expression gives rise to the enhanced grain number, with no impact on grain size.

### Spatial expression of GSK2 produces CL

Because spatial up-regulation of BRD3 in CL1 suppresses BR biosynthesis and BRs function through the BR signaling pathway, spatial suppression of BR signaling could also enable

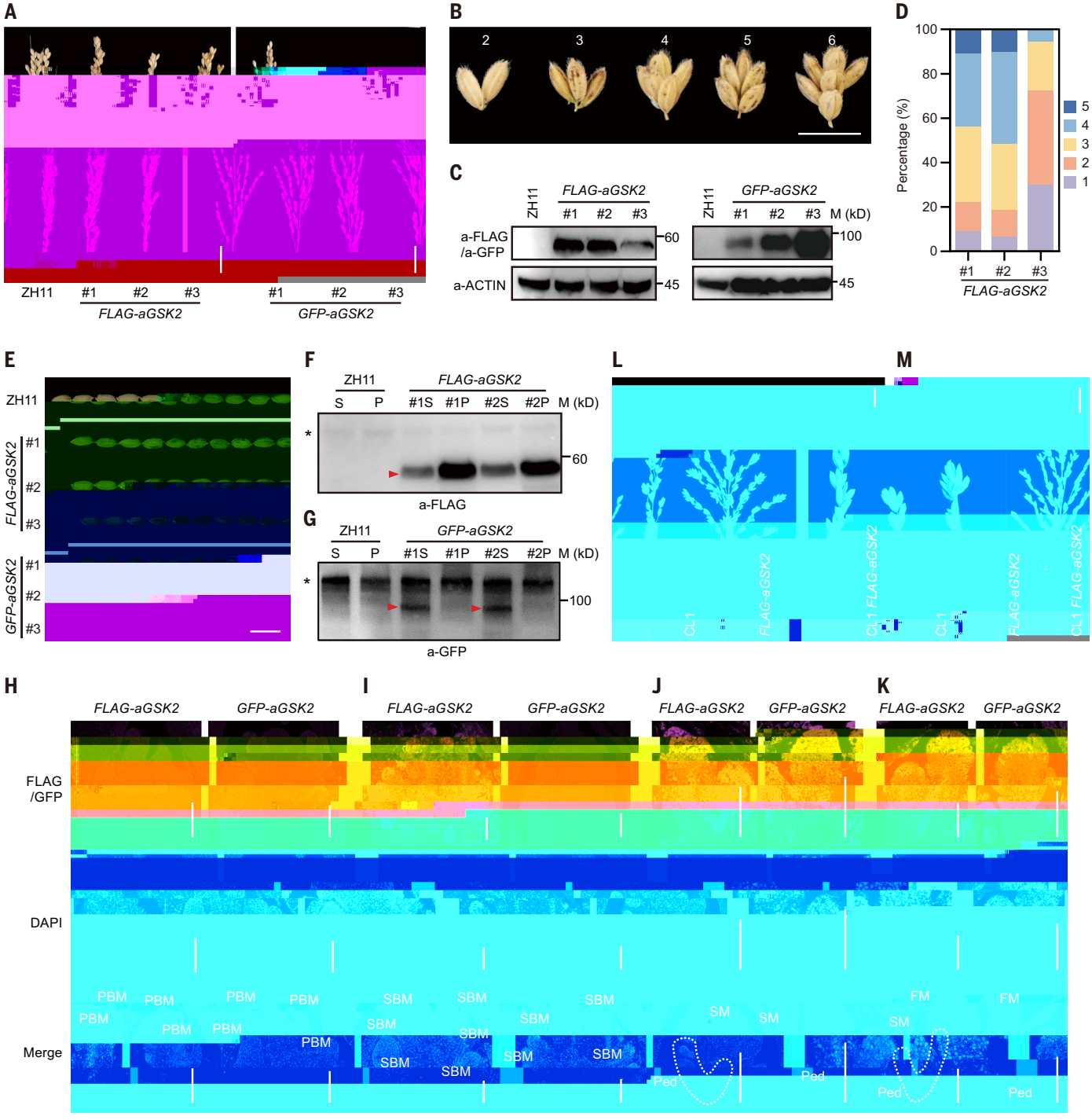
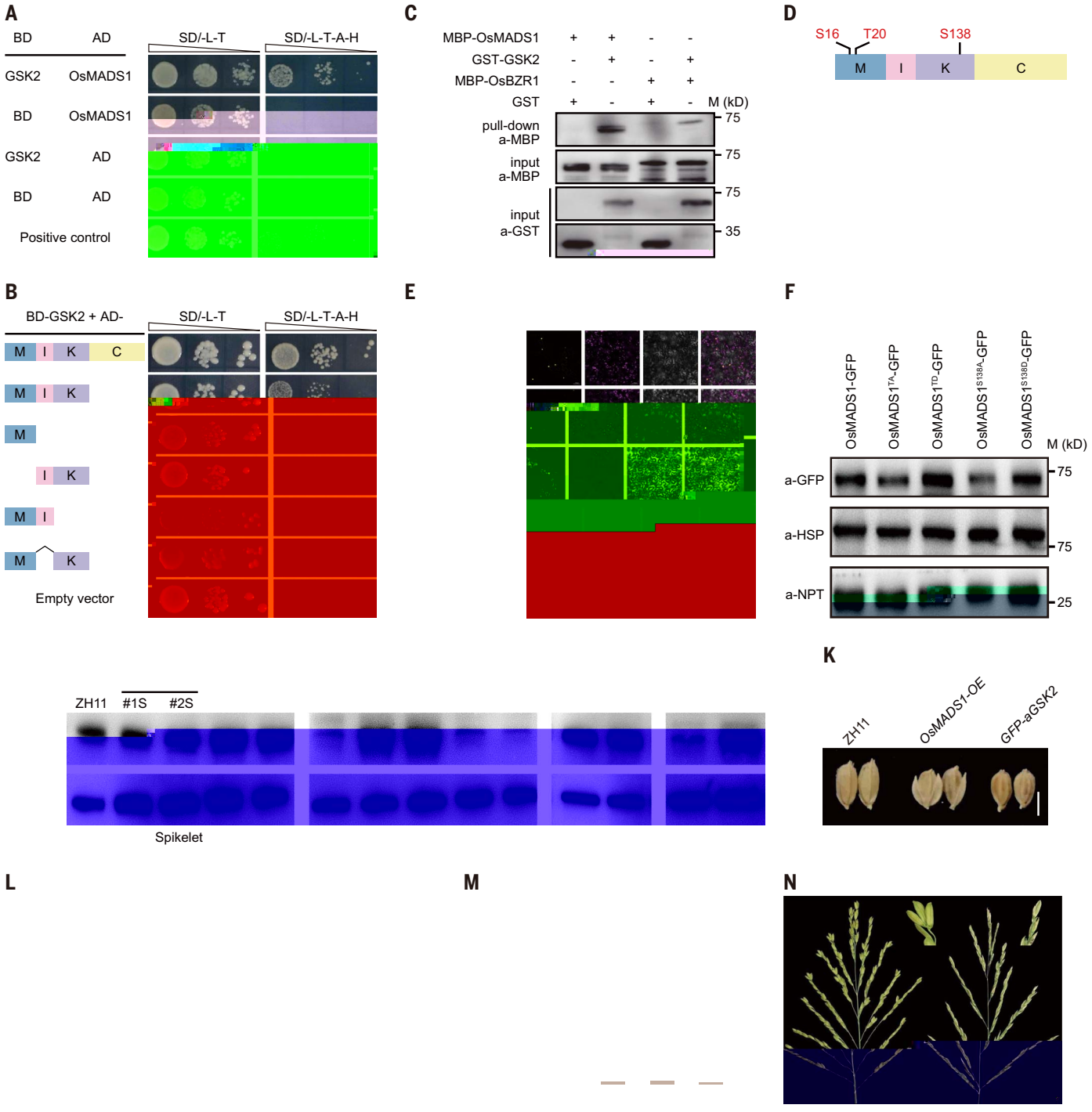



**Fig. 4. Spatially specific expression of GSK2 produces CL.** (**A**) Panicles of ZH11 and the transgenic plants. Scale bars: 2 cm. (**B**) Different cluster types in *FLAG-aGSK2* plants. Scale bar: 1 cm. (**C**) Detection of the fusion proteins in the transgenic plants by immunoblotting analyses. ACTIN was detected as control. M (kD), marker size indicated in kilodaltons. (**D**) Percentages of grain numbers belonging to different cluster types in the three *FLAG-aGSK2* lines, with 15 panicles analyzed for each line. (**E**) Comparison of grain size in different GSK2-related plants. Scale bar: 5 mm. (**F** and **G**) Detection of the fusion proteins in spikelets (S) and pedicels (P) of the plants by immunoblotting. Asterisk (*) represents a nonspecific band indicating equal protein loading. Arrowheads indicate target protein signals. (**H** to **K**) Immunofluorescence analysis of the fusion proteins during inflorescence development in *FLAG-aGSK2* and *GFP-aGSK2*. Ped, pedicel. The expression in the presumed pedicel positions were circled. Scale bars: 100 μm. (**L** and **M**) Comparison of the panicles and primary branches of CL1, *FLAG-aGSK2*, and the double mutant. Scale bars: 2 cm (L) and 1 cm (M).

OsMADS1-FLAG fusion protein (fig. S29). These results strongly suggested that phosphorylation of OsMADS1 by GSK2 enhances the stability of the OsMADS1 protein.

### OsMADS1 accumulates in pedicels of CLs

Given that GSK2 targets and promotes OsMADS1, the OsMADS1 protein should also specifically accumulate in CL plants in the same way that FLAG-aGSK2 protein or BRD3 transcripts do. To test this notion, we analyzed OsMADS1 protein levels in spikelets and pedicels of *FLAG-aGSK2* and *GFP-aGSK2* lines. Compared to wild

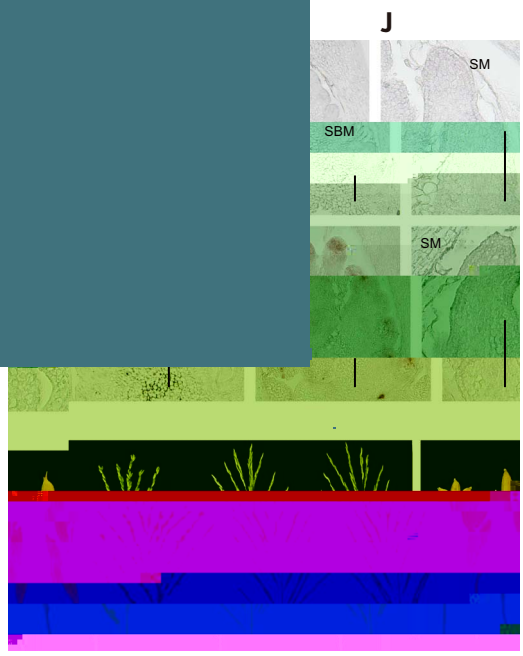

A
BD
AD
SD/-L-T
SD/-L-T-A-H
GSK2
OsMADS1
BD
OsMADS1
GSK2
AD
BD
AD
Positive control
B
BD-GSK2 + AD-
SD/-L-T
SD/-L-T-A-H
M
I
K
C
Empty vector
C
MBP-OsMADS1
GST-GSK2
MBP-OsBZR1
GST
M (kD)
pull-down
a-MBP
input
a-MBP
input
a-GST
75
75
75
35
D
S16
T20
S138
M
I
K
C
E
F
OsMADS1-GFP
OsMADS1TA-GFP
OsMADS1TD-GFP
OsMADS1S138A-GFP
OsMADS1S138D-GFP
M (kD)
a-GFP
a-HSP
a-NPT
75
75
25
ZH11
#1S
#2S
Spikelet
K
ZH11
OsMADS1-OE
GFP-aGSK2
L
M
N

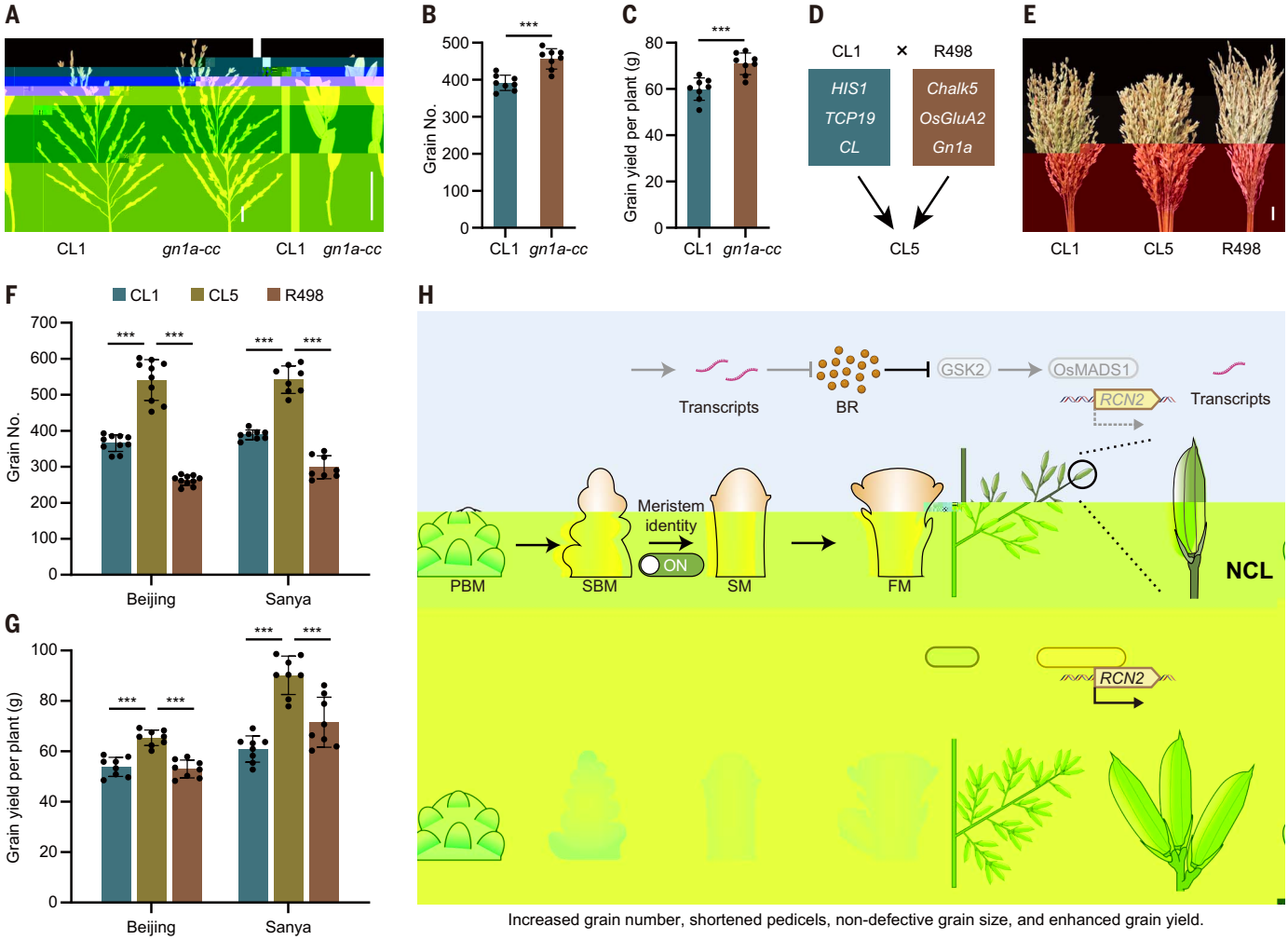
type, in spikelets, OsMADS1 was more abundant in *GFP-aGSK2*, but of similar abundance in *FLAG-aGSK2* (Fig. 5G). By contrast, in pedicels, OsMADS1 strongly accumulated in *FLAG-aGSK2*, but not in *GFP-aGSK2* (Fig. 5H). These data clearly demonstrated that the distribution patterns of OsMADS1 and GSK2 are highly similar in either *GFP-aGSK2* or *FLAG-aGSK2* (Figs. 4, F and G, and 5, G and H). In addition, the distribution patterns are also consistent with the plant phenotypes: *GFP-aGSK2* has smaller grains, whereas *FLAG-aGSK2* has shortened pedicels and normal grain size (Fig. 4E).

We then extended the analysis to CL1 and NCL1. Consistently, OsMADS1 levels were higher in pedicels, but not spikelets, of CL1 compared to NCL1 (Fig. 5, I and J). These patterns were similar to those observed in *FLAG-aGSK2*. Taken together, these results demonstrated that OsMADS1 acts downstream of GSK2 to mediate BR regulation of CL development.

### Knockdown of *OsMADS1* abolishes CL phenotype

Similar to GSK2, OsMADS1 has been characterized as a negative regulator of grain size (*35*, *36*). Furthermore, OsMADS1 has been implicated in regulating meristem identity and potentially enhancing spikelet number (*37*–*39*). Examination of *OsMADS1*-overexpressing plants (


J
SM
SBM
SM


A
CL1
gn1a-cc
CL1
gn1a-cc
B
Grain No.
CL1
gn1a-cc
C
Grain yield per plant (g)
CL1
gn1a-cc
D
CL1
×
R498
HIS1
TCP19
CL
Chalk5
OsGluA2
Gn1a
CL5
E
CL1
CL5
R498
F
CL1
CL5
R498
Grain No.
Beijing
Sanya
G
Grain yield per plant (g)
Beijing
Sanya
H
Transcripts
BR
GSK2
OsMADS1
RCN2
Transcripts
Meristem identity
ON
PBM
SBM
SM
FM
NCL
RCN2
Increased grain number, shortened pedicels, non-defective grain size, and enhanced grain yield.
I
TY2
TY2CL
J
Flower No. per node
Pedicel length (cm)
TY2
TY2CL
K
Castasterone (ng g−1)
6-Deoxocastasterone (ng g−1)
TY2
TY2CL
L
Rosa 'Parkdirektor Riggers'
Rosa 'Angela'
M
Flower No. per node
Pedicel length (cm)
PR
Angela
N
Castasterone (ng g−1)
6-Deoxocastasterone (ng g−1)
PR
Angela

number with 81.5 and 39.7% more grains per panicle compared to R498 and CL1, respectively (Fig. 7, E and F). In addition, CL5 possessed other advantageous traits such as a thick culm, heavier grains (fig. S35), and high yield (Fig. 7G), illustrating the promising potential of combining *CL* and *Gn1a* for enhancing rice yield.

### BR alteration in clustered growth in other species

Although we demonstrated that BR depletion controls CL in rice (Fig. 7H), we wanted to know whether this represent a general mechanism underlying clustered growth across plant species. Cluster peppers (*Capsicum annuum* L. var. *fasciculatum*) display multiple flowers clustered on a single flower bud formation node, whereas common peppers (*Capsicum annuum* L.) have only one flower attached to the node. We measured the BR content in pedicels of Tianyu 2 (TY2), a common pepper, and $TY2^{CL}$, a cluster pepper showing clustered flowers and shortened pedicels while closely resembling TY2 in vegetative and reproductive morphology (Fig. 7, I and J). Like CL1 and NCL1, the castasterone level was lower in $TY2^{CL}$ pedicels compared to TY2 (Fig. 7K). Conversely, the 6-deoxocastasterone level, the precursor of castasterone, was higher in $TY2^{CL}$ pedicels (Fig. 7K), indicating a possible loss of function in the BR synthase responsible for converting 6-deoxocastasterone to castasterone.

We also measured the BR content in the pedicels of the vine *Rosa chinensis* (*Rosa* "Parkdirektor Riggers") and the vine *Rosa* sp. (*Rosa* "Angela"), which displays the clustered flowers and shortened pedicels (Fig. 7, L and M). Like TY2 and $TY2^{CL}$, the castasterone level in *Rosa* "Angela" pedicels was lower compared to *Rosa* "Parkdirektor Riggers," whereas the 6-deoxocastasterone level exhibited the opposite trend (Fig. 7N). These consistent findings indicate that BR distribution may play a general role in regulating the clustered growth and inflorescence structure.

## Discussion

We unraveled the genetic basis underlying the historic CL germplasm, shedding light on the development of CL and its association with enhanced grain number, a crucial determinant of yield. We demonstrated that defective BR catabolism or signaling contributes to spikelet clustering and increased grain number, mediated by the specific spatial expression of *BRD3* transcripts, GSK2 protein, OsMADS1 protein, and *RCN2* transcripts. During CL's panicle development, the activation of *BRD3* in SBM triggers BR degradation, which enhances GSK2 stability, promotes OsMADS1 accumulation, and subsequently increases *RCN2* expression. As a result, the identity of SM is suppressed, leading to a delayed transition from BM to SM, ultimately yielding more SBMs and additional SMs formed adjacent to the terminal spikelet (Fig. 7H). Overall, our study exemplifies the successful utilization of natural rice germplasm resources harboring beneficial alleles to breed superior rice strains, from gene cloning to the dissection of the underlying molecular mechanisms.

KNOX transcription factors play a crucial role in the establishment and maintenance of the shoot apical meristem (SAM) (*48*). In *Arabidopsis*, BREVIPEDICELLUS (BP), a KNOX transcription factor, has been shown to regulate pedicel development (*48*). In rice, ORYZA SATIVA HOMEBOX1 (OSH1), the closest ortholog of BP, targets and activates BR catabolic genes, including *BRD3*, to maintain SAM activity (*49*). Therefore, the degradation of BRs should also promote the activity of SBM and SM. In our study, we observed an increase in grain number accompanied by shortened pedicels, and the expression of *BRD3* was detected sequentially in both the meristem cells and pedicel cells, suggesting a potential derivation of pedicels from meristem cells. Our findings imply that large structural variations preceding the *BRD3* gene may be responsible for the spatially specific increase in *BRD3* expression observed in CL. Although the detailed mechanism requires further exploration, one possibility is that these structural variations enhance chromatin accessibility, allowing the entry of transcription factors, such as OSH1, to promote *BRD3* expression.

Constitutive activation of *BRD3* in a transferred DNA (T-DNA) insertion mutant results in severe developmental defects, whereas moderately increased expression of *BRD3* in the heterozygote leads to higher grain number but smaller grains (*29*). Therefore, the practical application of *CL* in crop improvement can be attributed to two main factors related to *BRD3* expression: spatially specific expression and optimal expression. The resulting spatial distribution of BRs provides beneficial effects while preventing the reduction in grain size typically observed in BR-defective mutants, showcasing its practical applicability in crop improvement. Together with our previous research on the distribution of cytokinins and auxins (*50*, *51*), this work highlights the potential of optimizing hormone distributions to achieve desirable traits while mitigating negative associations in crop improvement efforts. Improving grain yield is intricately linked to expanding source supply, influenced by various factors such as source capacity, growth conditions (light, water, temperature, nutrients), and fertilizers. Although introduction of CL into several different modern cultivars all resulted in yield improvement, we observed distinct effects in different locations and genetic backgrounds. Considering that CL is influenced by BR content and each rice may exhibit variable BR levels, it is crucial to extensively test the effectiveness of CL in diverse contexts.

## Materials and methods summary

### Plant growth, suppressor screening, and segregating populations

For field plot analysis, each material was planted with a row interspace of 20 cm and plant interspace of 17.1 cm, with ~33 plants per square meter. At least four plots were planted as biological replicates. The border plants were removed from each plot to avoid margin effects during measurements. For experiments requiring seedling analysis, plants were grown in a growth chamber at 30°C with 10 hours light for day and 28°C with 14 hours dark for night. Half-strength Murashige and Skoog (½-MS) solution was provided as the nutrient source. For suppressor screening, dry CL1 seeds were incubated in sodium azide (1 mM in phosphate buffer pH 3.0) for 6 hours. After a thorough wash with tap water, the seeds were germinated and planted. A total of 10,000 individual lines were harvested, and 16 plants of each line were grown for mutant screening. Four segregating populations were generated by crossing CL1 with ZH11, 9311, *cl1-1*, and *cl1-2*, respectively. For each population, pooled DNAs from NCL plants and CL plants were prepared for BSA. Numbers of individual plants used for DNA pools from each population were as follows: 36 NCL and 34 CL from CL1 × ZH11, 24 NCL and 31 CL from CL1 × 9311, 31 NCL and 33 CL from CL1 × *cl1-1*, and 20 NCL and 19 CL from CL1 × *cl1-2*.

### RNA in situ hybridization

Specific fragments of *BRD3* and *RCN2* were amplified by PCR with primers listed in table S1. The fragments were subsequently inserted into the pEASY-Blunt Simple Cloning Vector (TransGen) for in vitro RNA transcription. Sense and antisense RNA probes were produced using T7 transcriptase and labeled with digoxigenin (Roche). Young panicle tissues from NCL1 and CL1 were collected and fixed in FAA (45% ethanol, 6% glacial acetic acid, and 5% formaldehyde) at 4°C overnight in a vacuum, dehydrated, and embedded in Paraplast Plus (Sigma-Aldrich, St. Louis). Tissue sections were prepared using a microtome (RM2235; Leica, Wetzlar) and then affixed to Poly-Prep slides (Sigma-Aldrich, St. Louis).

### In vitro phosphorylation and mass spectrum analysis

One microgram of GST-GSK2 and 2 μg of MBP-OsMADS1 or $MBP\text{-}OsMADS1^{TA}$ were incubated in the kinase buffer [10 mM Tris-HCl (pH 7.4), 50 mM NaCl, 6 mM $MgCl_2$, 0.5 mM dithiothreitol] with or without ATP and CIP (NEB) at 30°C for 2 hours. Phos-tag™ SDS–polyacrylamide gel electrophoresis (PAGE) was used to separate the phosphorylation protein following the instruction of the product Phos-tag™ Acrylamide (Wako). To identify the phosphorylation sites, 4 μg of recombinant

MBP-OsMADS1 or MBP-OsMADS1$^{TA}$ proteins were phosphorylated by 2 μg of GST-GSK2 for 1 hour in vitro. Phos-tag$^{TM}$ SDS-PAGE was used to separate the phosphorylation protein. The target protein bands were cut after one-step blue staining (Biotium). The LC-MS/MS detection and analysis were conducted by the Shanghai Luming Biological Technology Co., Ltd. (Shanghai, China).

### Immunofluorescence detection

An 8-μm-thick section was cut from paraffin-embedded young panicle tissues for detection. The slides were deparaffinized, rehydrated, and subjected to epitope retrieval by boiling in 1× All-purpose Powerful Antigen Retrieval Solution (Beyotime) for 20 min at 95°C, and set to cool for 30 to 40 min. The slides were washed two times in 1× Immunol Staining Wash Buffer (Beyotime) for 10 min each, blocked in Immunol Staining Blocking Buffer (Beyotime) for 60 min at room temperature, and then rinsed two times in 1× Immunol Staining Wash Buffer (Beyotime) for 10 min each. The primary antibody for OsMADS1 (1:50; Abclonal, Cat#A20328), or for FLAG-aGSK2 (anti-FLAG, 1:50; Sigma, Cat#F1804), or for GFP-aGSK2 (anti-GFP, 1:50; Abmart, Cat#M2004l), was used for incubation overnight at 4°C. The specimen was rinsed three times in 1× Immunol Staining Wash Buffer (Beyotime) for 5 min each, and then incubated in the Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor$^{TM}$ Plus 555 (1:400; Invitrogen, Cat#A32732) or Goat anti-Mouse IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (1:400; Invitrogen, Cat#A32723) at room temperature for 60 min in dark. The slides were further rinsed three times in 1× Immunol Staining Wash Buffer (Beyotime) for 5 min each, and then covered using coverslips filled with Antifade Mounting Medium with 4′,6-diamidino-2-phenylindole (DAPI; Beyotime). Fluorescence was captured with a confocal laser scanning microscope (LSM 980; Zeiss, Oberkochen).

### ChIP-seq and RNA-seq

About 1 g of young panicles at the 5-mm stage of *OsMADS1-OE* were used for ChIP-seq. Tissue fixation, nuclei extraction, and chromatin immunoprecipitation were performed using anti-OsMADS1 (1:100; Abclonal, Cat#A20328) antibody. The ChIP-seq DNA libraries were sequenced using the Illumina HiSeqTM2000 platform. After quality control, BWA software was used to align the clean reads against the Nipponbare reference genome (IRGSP1.0). MACS software was used for peak calling on a genome-wide basis, and the threshold for screening significant peak was $q$-value < 0.05. Significant peaks were assigned to the nearest gene. For RNA-seq, about 1 g of young panicles at the 5-mm stage of *OsMADS1-OE* and ZH11 were collected, and total RNA was extracted from each of three biological replicates using TRIzol reagent (Invitrogen). Sequencing libraries were generated using NEBNext Ultra RNA Library Prep Kit for Illumina (NEB), following the manufacturer's recommendations, and sequenced on Illumina NovaSeq platform. After cleaning up raw sequence reads, the clean reads were mapped to the Nipponbare reference genome (IRGSP1.0). The differentially expressed genes were analyzed using the edgeR package. Genes with a false discovery rate (FDR) < 0.05 and $\log_2$(fold change) > 1 were assigned as differentially expressed.

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## SUPPLEMENTARY MATERIALS

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Materials and Methods
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MDAR Reproducibility Checklist

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