Genome-wide identification of grain filling genes regulated by the OsSMF1 transcription factor in rice

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Abstract

Background: Spatial-and temporal-specific expression patterns are primarily regulated at the transcriptional level by gene promoters. Therefore, it is important to identify the binding motifs of transcription factors to better understand the networks associated with embryogenesis.

Results: Here, we used a protein-binding microarray (PBM) to identify the binding motifs of OsSMF1, which is a basic leucine zipper transcription factor involved in the regulation of rice seed maturation. OsSMF1 (previously called RISBZ1 or OsbZIP58) is known to interact with GCN4 motifs (TGA(G/C)TCA) to regulate seed storage protein synthesis, and it functions as a key regulator of starch synthesis. Quadruple 9-mer-based PBM analysis and electrophoretic mobility shift assay revealed that OsSMF1 bound to the GCN4 (TGA(G/C)TCA), ACGT (CCACGT(C/G)), and ATGA (GGATGAC) motifs with three different affinities. We predicted 44 putative OsSMF1 target genes using data obtained from both the PBM and RiceArrayNet. Among these putative target genes, 18, 21, and 13 genes contained GCN4, ACGT, and ATGA motifs within their 1-kb promoter regions, respectively. Among them, six genes encoding major grain filling proteins and transcription factors were chosen to confirm the activation of their expression in vivo. OsSMF1 was shown to bind directly to the promoters of Os03g0168500 (GCN4 motif), patatin-like gene (GCN4 motif), α-globulin (ACGT motif), rice prolamin box-binding factor (RPBF) (ATGA motif), and ONAC024 (GCN4 and ACGT motifs) and to regulate their expression.

Conclusions: The results of this study suggest that OsSMF1 is one of the key transcription factors that functions in a wide range of seed developmental processes with different specific binding affinities for the three DNA-binding motifs.

Keywords: OsSMF1, DNA-binding motif, Transcription factor, Grain filling genes

Background

Transcription is known to be regulated by the binding of transcription factors to their cognate motifs in the promoter regions of genes. In monocotyledons, several cis-elements, such as the prolamin box (TGTAAG), GCN4 motif (TGA(G/C)TCA), and AACA motif (AACAAAA), are highly conserved in the promoters of seed storage protein (SSP)-encoding genes and play central roles in controlling endosperm-specific gene expression during seed maturation (Takaiwa et al. 1996; Wu et al. 2000). Maize opaque-2 (O2) is an endosperm-specific transcription factor belonging to the bZIP family that has been shown to bind to the ACGT motif of the maize 22-kDa zein promoter and activate transcription (Schmidt et al. 1992). In addition, a barley bZIP transcriptional activator (BLZ1) binds to the GCN4 motif, which is putatively involved in regulating gene expression in the endosperm (Vicente-Carbajosa et al. 1998). Further, rice endosperm bZIP, RISBZ2 (also called REB), and opaque-2 heterodimerizing protein 1 (OHP1) specifically bind to the GCCACGT(A/C)AG sequence in the α-globulin (α-Glb) gene promoter and the TCCACG-TAGA sequence in the 22-kDa zein promoter, respectively (Nakase et al. 1997; Pysh et al. 1993). Other transcription factors are also involved in regulating the

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expression of starch synthesis genes. For example, a MYC-like protein (OsBP-5) and OsEBP-89, a member of the ethylene-responsive element-binding protein (EREBP) family, act synergistically as a heterodimer to regulate transcription of the rice Wx gene (Zhu et al. 2003). Additionally, rice starch regulator 1 (RSR1), which is an EREBP-type transcription factor, negatively regulates starch biosynthesis, and an RSR1-deficient mutant has been shown to exhibit the enhanced expression of starch synthesis genes in seeds (Fu and Xue 2010).

OsSMF1 (previously called RISBZ1) is a basic leucine zipper transcription factor that is involved in the regulation of rice seed maturation. It belongs to the maize O2-like protein group and is known to interact with the GCN4 motif (TGA(G/C)TCA) to regulate SSPs (Onodera et al. 2001). A previous study has shown that OsSMF1 (alternatively named OsbZIP58) is also involved in seed development, including the regulation of starch biosynthesis genes (Wang et al. 2013). It specifically binds to the ACGT elements in the promoters of both granule-bound starch synthase (GBSS) and starch branching enzyme (SBE1) (Wang et al. 2013). OsSMF1 gene expression is restricted to seeds, where it precedes the expression of storage protein-encoding genes (Onodera et al. 2001). Studies have demonstrated that OsSMF1 is involved in the regulation of starch synthesis, in addition to that of SSP synthesis, in the endosperm (Onodera et al. 2001; Wang et al. 2013). Here, we aimed to identify target genes for further functional analysis of OsSMF1.

For the large-scale identification of transcription factor target sites, several laboratory techniques, such as chromatin immunoprecipitation (ChIP) and DNA adenine methyltransferase identification (DamID), have been devised (Orian 2006; Pokholok et al. 2002; Ren et al. 2000; van Steensel et al. 2001; Wyrick et al. 2001). Recently, chip-based protein-binding microarrays (PBMs) have been developed, allowing for the identification of protein-DNA interactions in vitro. In our previous study, we constructed a quadruple 9-mer protein-binding microarray (Q9-PBM), in which 131,072 quadruple probes were designed to cover all possible combinations of 9-mers of the reverse complementary sequences (Kim et al. 2009). The specificity of Q9-PBM was confirmed using well-known DNA-binding sequences, including Cbf1 and CBF1/DREB1B, and it was also used to elucidate the unidentified cis-acting element of the ONAC024 rice transcription factor (Kim et al. 2009). The Q9-PBM has mainly been used to assess the interactions of transcription factors with short synthetic DNA sequences and to evaluate their DNA sequence specificities. Here, we used the Q9-PBM to determine the binding motifs of OsSMF1. In a previous study, the RiceArrayNet (RAN) was developed from collective data obtained from 60 K rice microarrays (Lee et al. 2009), and it has been widely used (Hamada et al. 2011; Lorenz et al. 2011; Movahedi et al. 2011). We have constructed a new version of the RAN that provides co-expression information on genes, including correlation coefficients, calculated using accumulated data obtained from 300 K rice microarrays (http://bioinfo.mju.ac.kr/arraynet/rice300k_2011/query/).

To predict the co-expressed genes, we also examined the relationships among genes that are putatively regulated by OsSMF1 by RAN analysis.

Q9-PBM analysis revealed that OsSMF1 recognized three binding motifs, namely the GCN4 [TGA(G/C)TCA], ACGT [CCACGT(G/C)], and ATGA (GGATGAC) motifs, with different affinities. We detected 85 genes that were positively regulated by OsSMF1 using the RAN analysis under the tested conditions, with a minimum correlation value of 0.55 and a depth of 1. Among these putative target genes, 18 (21.2%), 21 (24.7%), and 13 (15.3%) contained GCN4, ACGT, and ATGA motifs within their 1-kb promoter regions, respectively. In addition to confirming the known OsSMF1 target genes, we predicted 35 potential target genes that have not been previously described in immature seeds. Using qRT-PCR and the protoplast transactivation assay, we found that in vivo, OsSMF1 activated Os03g0168500 and patatin-like protein, which contain the GCN4 motif; ONAC024 and ONAC026, which contain either the GCN4 or ACGT motifs; and prolamin box-binding factor (RPBF) and CCCH-type zinc finger protein (OsGZF1), which contain the ATGA motif. The results suggest that OsSMF1 has specific binding affinities for the three motifs and that it functions in a wide variety of seed developmental processes.

Results
Identification of the multiple binding motifs of OsSMF1 by PBM
OsSMF1 is a transcription factor previously known as RISBZ1 or OsbZIP58. In this study, OsSMF1 was mainly expressed in the endosperm at 11 and 21 DAF during seed maturation (Additional file 1: Figure S1) (Onodera et al. 2001; Wang et al. 2013). A previous classification of bZIP proteins from green plants resulted in the identification of five rice bZIP transcription factors, including OsSMF1, belonging to the same group (Correa et al. 2008). In this study, sequence analyses were performed using ClustalW, and the results revealed that OsSMF1 possessed a high degree of similarity of 52.7% with the REM (RISBZ2) protein, whereas it shared approximately 31% sequence identity with the RITA-1 (RISBZ3), RISBZ4, and RISBZ5 proteins (Fig. 1a). Additionally, it possessed approximately 51.3 and 49.6% amino acid sequence identities with maize OHP1 and barley BLZ1, respectively, which contained conserved sequences
within the leucine zipper regions with homologies of 73.7–76.3% (Fig. 1b). These endosperm-specific transcription factors with similarities to OsSMF1 are known to bind to either the GCN4 or ACGT motif.

We first attempted to identify the binding motifs of the OsSMF1 protein using a Q9-PBM. This transcription factor, as well as similar variants, bind to multiple binding motifs (Godoy et al. 2011). To this end, the full-length OsSMF1 cDNA was fused at the N-terminus to the DsRed fluorescent protein. Purified recombinant OsSMF1-DsRed protein expressed in E. coli was hybridized to the Q9-PBM. Then, the consensus binding motifs were determined based on signal strength (Jung et al. 2012; Kim et al. 2009).

The signal distribution curve of the OsSMF1 PBM was characterized by a deep leftward slope, followed by a right tail (Additional file 2: Figure S2); the shape of the curve was attributed to specific interactions between the protein and features on the microarray. For motif extractions, 1,286 total signals in the steep left region with high intensities were clustered and identified using SEQLOGO (Fig. 2a, c and e). We found significant binding motifs for OsSMF1, including ACGT (CCACGTCA), with a high intensity of 13,715, GCN4 (TGAGTCA) with a moderate intensity of 7,639, and ATGA (GGATGAC) with an intensity of 6,463 (Fig. 2a, c and e). In addition, we found that OsSMF1 bound to a new motif termed the ATGA motif, which can be considered a novel OsSMF1 binding motif. To determine the flanking sequence of the ACGT core sequence that is essential for DNA binding by OsSMF1, we analyzed the relative signal intensities of single nucleotide substitution variants of the putative binding sequences (Fig. 2b). In agreement with the Q9-PBM results, individual substitutions at all positions of the CCACGTC sequence resulted in significantly reduced binding signal intensities. Although C had a stronger binding affinity than G at the seventh nucleotide in the ACGT motif, G also had a relatively strong affinity for OsSMF1. Thus, we found a significantly high affinity of OsSMF1 for the ACGT motif, CCACGT(C/G), which may be considered an OsSMF1-specific binding motif (Fig. 2).

To further analyze the multiple binding motifs of OsSMF1 with different intensities, we assayed their binding specificities to recombinant OsSMF1 by electrophoretic mobility shift assay (EMSA) using biotinylated...
double-stranded oligonucleotide probes corresponding to each motif. The binding of OsSMF1 to the 21-bp fragments of the GCN4, ACGT, and ATGA motifs was detected as lagging bands (Fig. 3). As shown in Fig. 3a, c and e, three bases of the 21-bp motifs were sequentially mutagenized, and these mutants were used as competitors (100-fold increased molarities) in the EMSAs. The results revealed that the introduction of mutagenized nucleotides into any part of the core motifs as competitors had little or no effect on the binding of native fragments, whereas mutations flanking the core motifs led to the loss of binding. Taken together, these results suggested that OsSMF1 specifically bound to the GCN4, ACGT, and ATGA motifs with high affinity (Fig. 3a, c and e). For the determination of binding affinities, dissociation constants ($K_d$) were estimated by gel shift assay using SYBR Gold. DNA-binding band intensity was measured at various DNA substrate concentrations (0, 0.1, 0.2, 0.4, 0.8, 1.2, 2, 4, and 6 μM) in the presence of OsSMF1 (0.2 μg/μl). The $K_d$ values for the binding of OsSMF1 to the GCN4, ACGT, and ATGA motifs were 0.6458 μM, 0.3353 μM, and 1.117 μM, respectively (Fig. 3b, d and f). The binding affinities of OsSMF1 for the three DNA-binding motifs, as indicated by the $K_d$ values, showed the same trends as those revealed by the Q9-PBM, indicating that OsSMF1 was able to bind to the GCN4, AGCT, and ATGA motifs independently with different affinities.

**Gene expression network of OsSMF1 based on microarray expression data**

OsSMF1 is considered an important transcription factor involved in the regulation of SSPs and starch synthesis. To further elucidate the regulatory mechanisms of OsSMF1 in the cereal endosperm, we predicted its target genes. In a previous paper, the RAN was developed using accumulated data from 60 K rice microarrays (Lee et al. 2009), and the new version of the RAN has been used to elucidate gene functions, as it provides co-expression information for genes, including correlation coefficients, obtained from 174 300 K microarrays of the rice genome (*Oryza sativa*) (http://bioinfo.mju.ac.kr/arraynet/rice300k_2011/) (Hamada et al. 2011; Lorenz et al. 2011; Movahedi et al. 2011). We identified 85 genes that were positively correlated with OsSMF1 in rice using the RAN under the tested conditions, with a minimum correlation value of 0.55 and a depth of 1 (Additional file 3: Table S2, Additional file 4: Figure S3). Among them, 21 (24.7%), 18 (21.2%), and 13 (15.3%) putative target genes contained ACGT, GCN4, and ATGA motifs in their 1-kb promoter regions, respectively (Table 1).

These predicted target genes were functionally categorized into 4 groups according to the Gene Ontology analysis (http://geneontology.org/) (Table 2). Among the 44 predicted genes, 6 were categorized under the term...
nutrient reservoir activity (GO:0045735). Among these 6 genes, five glutelin genes (GluA-1, GluA-2, GluA-3, GluB-4, and GluB-5) contained the GCN4 motif with or without the other motifs, and one globulin gene (α-glub) contained the ACGT motif. Previous reports (Kawakatsu et al. 2008; Yamamoto et al. 2006) have indicated that OsSMF1 initiates trans-activation from the promoters of the GluA-1, GluA-2, and GluA-3 genes following recognition of the GCN4 motif. GBSS, a known target gene of OsSMF1, was also identified in our analysis (Wang et al. 2013). Our results consistently demonstrate the involvement of OsSMF1 not only in starch synthesis but also in the regulation of SSPs in developing seeds. Five and three additional genes were functionally categorized under the terms “defense response (GO:0006952)” and “negative regulation of RNA metabolic process”

Fig. 3 Identification of OsSMF1 binding motifs in vitro. a, c, e EMSA-based competition analysis of OsSMF1. Sequences of oligonucleotides used as probes and competitors are depicted. Wild type, 21-bp sequences containing the GCN4 (a), ACGT (c), and ATGA motifs (e); and M1 to M6, 21-bp sequences with three nucleotide mutations. The GCN4, ACGT, and ATGA motifs are underlined. The OsSMF1-DsRed fusion protein was used for EMSA with the 21-bp sequences. Competitors were added in 100-fold molar excess. Lane 1, no protein; Lane 2, no competitor; and lanes 3 to 9, with competitors. b, d, f Accurate determination of the dissociation constant $K_d$ by gel shift assays using SYBR Gold. The DNA-binding band intensities determined using Gel-Pro Analyzer program were plotted against the various DNA substrate concentrations. The measured $K_d$ values were $0.6458 \mu M$ for the binding of OsSMF1 to the GCN4 motif (b), $0.3353 \mu M$ for ACGT (d), and $1.117 \mu M$ for ATGA (f).
| IRGSP Gene | r-value | Motif | GCN4 | ACGT | ATGA | Description |
|------------|---------|-------|------|------|------|-------------|
| Os01g0762500 | 0.63 | 889 | tga-gtc | GluA-1 |
| Os10g0400200 | 0.56 | 897 | tga-gtc | GluA-2 |
| Os01g0898500 | 0.64 | 773 | tga-gtc | Patatin-like protein 2 (PAT) |
| Os06g0133000 | 0.55 | 185 | tga-gtc | Granule-bound starch synthase I, chloroplast precursor (GBSS) |
| Os03g0168500 | 0.63 | 496 | tga-gtc | Endosperm-specific gene 44 (OsEnS44) |
| Os03g0712800 | 0.63 | 200 | tga-gtc | Glutamine synthetase root isozyme 2 |
| Os01g0393400 | 0.62 | 572 | tga-gtc | MDR-like ABC transporter |
| Os03g0799600 | 0.59 | 702 | tga-gtc | ES43-like protein |
| Os04g0437000 | 0.58 | 512 | tga-gtc | No apical meristem (NAM) domain-containing protein (ONAC79) |
| Os02g0459400 | 0.55 | 272 | tga-gtc | Conserved hypothetical protein |
| Os03g0179700 | 0.54 | 932 | tga-gtc | Protein of unknown function DUF567 family protein |
| Os03g0427300 | 0.66 | 898 | 381 | ccacgtc | GluA-3 |
| Os05g0415400 | 0.55 | 850 | 352 | ccacgtg | No apical meristem (NAM) domain-containing protein (ONAC024) |
| Os04g0528000 | 0.66 | 716 | 366 | ccacgtg | Protein of unknown function DUF789 family protein |
| Os02g0268100 | 0.59 | 832 | tga-gtc | 171 | ggatgac | GsB-5 |
| Os02g0268300 | 0.57 | 818 | tga-gtc | 185 | ggatgac | GsB-4 |
| Os09g0520400 | 0.63 | 715 | tga-gtc | 744 | ggatgac | OsbZIP76 |
| Os02g0249500 | 0.61 | 105 | tga-gtc | 802 | ggatgac | Hypothetical protein |
| Os05g0499100 | 0.55 | 436 | ccacgtc | 26-kDa globulin (alpha-globulin) |
| Os07g0182000 | 1.00 | 841 | ccacgtc | OsSMF1 |
| Os11g0582400 | 0.56 | 100 | ccacgtc | Endosperm-specific protein (OsEnS146) |
| Os01g0393100 | 0.60 | 257 | ccacgtc | No apical meristem (NAM) domain-containing protein (ONAC026) |
| Os05g0569400 | 0.63 | 650 | ccacgtg | Galactose mutarotase-like domain-containing protein. |
| Os01g0133400 | 0.61 | 381 | ccacgtg | Hexose transporter (fragment) |
| Os02g0694100 | 0.59 | 534 | ccacgtc | Cyclin-like F-box domain-containing protein |
| Os03g0598700 | 0.59 | 32 | ccacgtc | Zn-finger, RING domain-containing protein |
| Os05g0404700 | 0.59 | 342 | ccacgtc | Methyl-CpG-binding protein (MBD1) |
| Os11g0701100 | 0.59 | 832 | ccacgtg | Xylanase inhibitor protein 2 |
| Os01g0585100 | 0.58 | 966 | ccacgtc | Conserved hypothetical protein |
| Os06g0141400 | 0.57 | 10 | ccacgtc | Early nodulin |
| Os05g0557600 | 0.57 | 226 | ccacgtc | Ubiquitin-conjugating enzyme domain-containing protein |
| Os04g0508500 | 0.56 | 397 | ccacgtc | Snapdragon myb protein 305 homolog |
| Os05g0491400 | 0.56 | 196 | ccacgtc | LRR protein |
| Os01g0517800 | 0.56 | 862 | ccacgtg | PLATZ transcription factor |
| Os04g0629300 | 0.55 | 263 | ccacgtc | SNF2-related domain-containing protein |
| Os10g0138600 | 0.69 | 240 | ccacgtc | 158 | ggatgac | Cyclin-like F-box domain-containing protein |
Further, 10 transcription factors, including OsSMF1, were assigned to "regulation of nitrogen compound metabolic process (GO:0051171)." Among these transcription factors, two, including OsGZF1 and rice prolamin box-binding factor (RPBF), were co-expressed with OsSMF1 (Chen et al. 2014; Yamamoto et al. 2006). These in vitro results suggest that the OsSMF1 gene may be involved in the regulation of a wide range of target genes involved in seed development during embryogenesis.

### Table 1: Candidate OsSMF1 target genes, the binding motifs and RAN analysis and microarray expression data (Continued)

| Accession | Name | IRGSP Gene | GCN4 | ACGT | ATGA | Description |
|-----------|------|------------|------|------|------|-------------|
| Os12g0151000 | 0.69 | ccacgtg 426 | 233 ggtgcc | Purple acid phosphatase |
| Os02g0252400 | 0.66 | ccacgtg 737 | ggtgcc | Rice prolamin box-binding factor (RPBF) |
| Os07g0668600 | 0.65 | ccacgtg 360 | ggtgcc | Zinc finger, CCCH-type domain-containing protein (OsGZF1) |
| Os04g0677400 | 0.66 | ccacgtg 975 | ggtgcc | Conserved hypothetical protein |
| Os07g0457400 | 0.62 | ccacgtg 764 | ggtgcc | Conserved hypothetical protein |
| Os03g0766200 | 0.61 | ccacgtg 770 | ggtgcc | Bifunctional inhibitor/plant lipid transfer protein/seed storage domain-containing protein |
| Os07g0485100 | 0.60 | ccacgtg 589 | ggtgcc | Nitrilase/cyanide hydratase and apolipoprotein N-acyltransferase domain-containing protein |
| Os07g0603600 | 0.56 | ccacgtg 237 | ggtgcc | Alpha/beta hydrolase family protein |

### Table 2: Gene Ontology classification of putative OsSMF1 target genes

| Accession | Name | Ontology | IRGSP Gene | Description |
|-----------|------|----------|------------|-------------|
| GO0045735 | nutrient reservoir activity | molecular_function | Os01g0762500 | GluA-1 |
| | | | Os10g0400200 | GluA-2 |
| | | | Os03g0427300 | GluA-3 |
| | | | Os02g0268300 | GluB-4 |
| | | | Os02g0268100 | GluB-5 |
| | | | Os05g0499100 | 26-kDa globulin (alpha-globulin) |
| GO0051171 | regulation of nitrogen compound metabolic process | biological_process | Os02g0268100 | OsSMF1 |
| | | | Os01g0393100 | No apical meristem (NAM) domain-containing protein (ONAC026) |
| | | | Os05g0404700 | Methyl-CpG-binding protein (MBD1) |
| | | | Os04g0629300 | SNF2-related domain-containing protein. |
| | | | Os05g0415400 | No apical meristem (NAM) domain-containing protein (ONAC024) |
| | | | Os03g0799600 | ES43-like protein |
| | | | Os04g0437000 | No apical meristem (NAM) domain-containing protein (ONAC79) |
| | | | Os09g0520400 | OsbZIP76 |
| | | | Os02g0252400 | Rice prolamin box-binding factor (RPBF) |
| | | | Os07g0668600 | CCCH-type Zinc finger protein (OsGZF1) |
| GO0006952 | defense response | biological_process | Os11g0582400 | Endosperm-specific protein (OsEnS146) |
| | | | Os11g0701100 | Xylanase inhibitor protein 2 |
| | | | Os03g0197900 | Protein of unknown function DUF567 family protein |
| | | | Os02g0268100 | GluB-5 |
| | | | Os02g0268300 | GluB-4 |
| GO0051253 | negative regulation of RNA metabolic process | biological_process | Os05g0404700 | Methyl-CpG-binding protein (MBD1) |
| | | | Os03g0799600 | ES43-like protein |
| | | | Os04g0629300 | SNF2-related domain-containing protein |
Expression of target genes enhanced by overexpression of OsSMF1
To assess whether the putative target genes described above are biologically significant in vivo, a vector was constructed in which OsSMF1 was expressed under control of the promoter of Wsi (a member of the group 3 Lea family), which is predominantly active in the whole grain, including the endosperm, embryo, and aleurone layer, during seed development (Yi et al. 2011). Microarray analysis further showed high Wsi mRNA expression in the callus (Additional file 1: Figure S1). The bialaphos resistance (bar) gene was used as a selectable marker to identify transgenic calli. Four independent transgenic plants were obtained using the Agrobacterium-mediated transformation method (Sohn et al. 2006). Calli were generated from T1 seeds, and the expression of OsSMF1 and its target genes was measured by RT-PCR and qRT-PCR.

The overexpression of OsSMF1 was confirmed by RT-PCR and qRT-PCR, which demonstrated that expression of this gene was upregulated by 200- to 1000-fold in calli derived from OsSMF1-transformed plants compared with that in calli derived from wild-type and nullizygous plants (Figs. 4 and 5). We also confirmed that the expression of α-Glb and GBSS, which are known target genes of OsSMF1 but were not expressed in the calli, was increased in OsSMF1-transformed calli compared with that in nullizygous and wild-type calli, as shown by RT-PCR (Fig. 4). These results revealed that OsSMF1 activated expression of the α-Glb and GBSS genes in vivo.

The expression of Os03g0168500 (endosperm-specific gene 44, OsEnS44) was markedly upregulated in four independent calli derived from OsSMF1-transformed plants compared with that in nullizygous and wild-type calli (Fig. 5). The promoter region of OsEnS44 contains GCN4 motifs in its 1-kb promoter region. Os11g0582400 (endosperm-specific protein 146, OsEnS146) expression was also significantly increased in two independent OsSMF1-transformed calli (Fig. 5), and its promoter region contains the ACGT motif. Thus, our results indicated that OsEnS44 and OsEnS146 acted as novel OsSMF1 target genes in vivo. Two transcription factors identified as putative OsSMF1 target genes, RPBF and ONAC024, were selected for validation of their expression in the OsSMF1-transformed calli by qPCR. The promoter region of ONAC024 contains both the GCN4 and ACGT motifs, and the ATGA motif has been detected in the RPBF promoter. RPBF expression was increased by up to 4-fold in three independent OsSMF1-transformed calli compared with that in wild-type calli (Fig. 5). Further, ONAC024 expression was upregulated by approximately 2-fold in two independent OsSMF1-transformed calli (Fig. 5). These results also indicated that RPBF and ONAC024 could be considered target genes of OsSMF1. Previous studies have suggested that

![Fig. 4 Production of DsRed-fused OsSMF1-transformed rice.](image-url)

**Fig. 4** Production of DsRed-fused OsSMF1-transformed rice. **a** Schematic representation of the expression vector Wsi:SMF1 used for rice transformation. **b** The expression of OsSMF1 and its target genes was determined by RT-PCR analysis. Specifically, the expression of granule-bound starch synthase (GBSS) and 26-kDa globulin (α-Glb) were analyzed. Total RNA was extracted from calli. Tubulin expression is shown as a constitutive control.
RPBF and OsSMF1 (RISBZ1) synergistically activate seed-specific genes during grain filling (Kawakatsu et al. 2009; Yamamoto et al. 2006).

**Binding analysis of promoter regions targeted by OsSMF1**

To investigate whether OsSMF1 activates expression of the putative target genes listed in Table 2, we performed a transient activation assay in rice protoplasts using 13 putative genes: α-Glb, patatin-like protein, prolamin10.2, OsEnS44, OsEnS146, ONAC024, ONAC026, OsSMF1, aldose 1-epimerase family protein, methyl-CpG-binding protein (MBD1), PLATZ, RPBF, and OsGZF1. Their promoters were fused to fLUC (promoter:fLUC) as a reporter. Among the 13 putative genes, six genes were upregulated and one gene, OsGZF1, was downregulated by OsSMF1. The expression of OsEnS44 and OsEnS146, which contain the GCN4 and ACGT motifs in their promoter region, respectively, was dramatically increased in OsSMF1-transformed rice. RPBF and ONAC024 transcription factors, the promoters of which contain the ATGA and GCN4/ACGT motifs, respectively, were highly expressed in OsSMF1-transformed calli compared with wild-type calli. We also selected patatin-like protein, the promoter of which contains the GCN4 motif (Table 1), as a storage protein. The other selected genes, including patatin-like protein, showed similar expression patterns in both transgenic and wild-type calli (data not shown). The α-Glb promoter was used as a positive control. The Os12g0621600 (hydroxyproline-rich glycoprotein, HRGP) promoter, which was determined to be correlated with OsSMF1 in RAN analysis (correlation value of 0.6) but does not contain any of the three binding motifs in its promoter, was selected as a negative control (HRGP:fLUC). In these assays, the effector gene, OsSMF1, was expressed under control of the P35S promoter and co-transfected together with the reporter construct into the protoplast. Analysis of LUC activity demonstrated that the co-transfection of EnS44:fLUC, α-Glb:fLUC, and ONAC024:fLUC with OsSMF1 resulted in induction of the expression of the luciferase reporter gene by 23.7-, 33-, and 51.3-fold, respectively, compared with co-transfection of HRGP:fLUC (Fig. 6). The activity of fLUC by the promoters of the Patatin-like protein, ONAC026, and RPBF were increased up to 5.5-, 1.8-, and 2.2-fold in the presence of OsSMF1, but the promoter of OsGZF1 was downregulated by this transcription factor (Fig. 6). These results indicated that OsSMF1 regulated the expression of OsEnS44 and patatin-like genes by binding to the GCN4 motifs in their promoters. The expression of RPBF and ONAC026 was also directly regulated by the binding of OsSMF1 to the ATGA and ACGT motifs in the promoter, respectively. The expression of ONAC024, which contained both the GCN4 and ACGT motifs in its promoter, was greatly increased by OsSMF1 (Fig. 6).

**Discussion**

Determination of the DNA-binding specificities of transcription factors is important for the understanding of transcriptional regulatory codes. Rice seed development is directly related to crop yield and is finely controlled by complex regulatory networks. Gene co-expression network analysis has shown that transcription factors are involved in the complex regulation of rice seed development (Xue et al. 2012). However, many of the primary target genes of these transcription factors as part of regulatory networks remain to be elucidated. We aimed to identify target genes of the seed-specific transcription factor OsSMF1, which is known to be involved in the regulation of rice SSP gene expression (Kawakatsu
et al. 2009; Yamamoto et al. 2006) and starch synthesis (Onodera et al. 2001; Wang et al. 2013). Q9-PBM analysis, which is a powerful and rapid method for the identification of putative functional cis-regulatory elements, allows for the accurate quantification of the binding affinities of some transcription factor binding motifs using all possible 9-mer combinations of probes (Kim et al. 2009). We used this technology to characterize the binding specificities of OsSMF1 and identified three DNA-binding motifs for this transcription factor, including the GCN4 (TGA(G/C)TCA), ACGT (CCACGT(C/G)), and ATGA (GGATGAC) motifs (Figs. 2 and 3). It has recently become apparent that a transcription factor may interact with more multiple DNA elements. For example, *Arabidopsis* MYC2 binds with high affinity to the G-box (CACGTG), T/G (AACGTG), and G-like (CATGTG) motifs (Godoy et al. 2011). Another rice bZIP protein, REM, binds to both the ACGT and GCN4 motifs (Nakase et al. 1997). OsSMF1 and REM have a close phylogenetic relationship (52.9%). However, OsSMF1 expression is restricted to the mature seed, whereas REM is expressed in all tissues. Furthermore, the expression of OsSMF1 at 11 DAP was determined to be 6-fold higher than that of REM in this study (Additional file 1: Figure S1). These results suggest that OsSMF1 plays important roles in a wide range of biosynthetic processes during seed maturation. In addition to the known GCN4 binding motif, OsSMF1 also recognizes the ATGA and ACGT motifs. The ACGT motif has been previously shown to be a target of OsSMF1-regulated gene expression, but the essential flanking sequence of this motif has not yet been identified (Wang et al. 2013). Q9-PBM analysis and EMSA performed in this study revealed that CCACGT(C/G) is an essential sequence that is bound by OsSMF1 (Figs. 2 and 3). We found that OsSMF1 bound to TGAGTCA (GCN4 motif), CCACGT (ACGT motif), and GGATGAC (ATGA motif) with intensities of 6,463, 13,715, and 7,639, respectively. The $K_d$ value for the binding of OsSMF1 to the ACGT motif was approximately two- and four-fold higher than those for the GCN4 and ATGA motifs, respectively (Fig. 3), demonstrating that OsSMF1 plays prominent roles during seed development by binding to these three motifs in vitro.

First, we selected genes containing the GCN4, ACGT, and ATGA motifs in their 1-kb promoter regions from the RAP2 rice database. Second, the RAN analysis, which was based on the expression pattern correlation, was capable of narrowing down the putative target genes of OsSMF1 to a subset of 85 genes (Additional file 2: Figure S2). To validate these predicted genes, we searched the GCN4 motif for known OsSMF1 binding sequences by combined microarray analysis. Among the six putative genes, four genes (*GluA-1*, *GluA-2*, *GluA-3*, and *α-Glb*) have been previously identified as OsSMF1 target genes by transient assay using rice callus protoplasts (Yamamoto et al. 2006). Among the remaining targets that were detected, two globulin genes (*GluB-4* and *GluB-5*) were characterized as SSPs. A previous study demonstrated that OsSMF1 causes reduced activation of the *α-Glb* promoter compared with that of the *GluA-1* and *GluA-3* promoters (Yamamoto et al. 2006).
However, this group assessed the α-Glb region from −340 bp to +73 bp, which does not include the ACGT motif positioned at −436 bp relative to the ATG start codon, in transient assays using rice callus protoplasts. Although patatin-like protein was not classified under the Gene Ontology term “nutrient reservoir activity (GO:0045735),” it contained the GCN4 motif and was determined to be one of the direct target genes of OsSMF1 (Table 1, Fig. 6). Patatin represents approximately 40% of the soluble protein in potato tubers, but it has not yet been studied in rice.

Among the 44 total putative target genes, 9 transcription factors (27.5%), including OsSMF1, were observed, which were assigned to “regulation of nitrogen compound metabolic process (GO:0051171, Table 2). OsSMF1 is specifically expressed in the aleurone and subaleurone layers of the developing endosperm (Onodera et al. 2001), and the expression of the target genes of this transcriptional activator may be restricted to these subcellular locations. Among the 9 transcription factors, 6 genes, which were preferentially expressed at 11 and 21 DAP when OsSMF1 was also highly expressed (Additional file 5: Figure S4), were selected and four transcription factors were identified as target genes through in vivo protoplast analysis. The promoters of OsSMF1, ONAC026, and MBD1 contained the ACGT motif, and the OsGZF1 and RPBF genes incorporated the ATGA motif in their promoters. The ONAC024 promoter was selected, which contained both the GCN4 and ACGT motifs on its promoter, showed significantly increased expression by OsSMF1. ZmaNAC36, which was cloned from maize by homologous cloning using rice ONAC026, is co-expressed with many starch synthesizing genes. Thus, ONAC026 might be co-expressed with starch biosynthetic genes in the rice endosperm. A previous study showed that OsSMF1 (alternatively named OsbZIP58) null mutants displayed abnormal seed morphology with altered starch accumulation and decreased amounts of total starch, particularly amylase, with binding to the promoters of six starch-synthesizing genes (Wang et al. 2013). We suggest that ONAC024 and ONAC026 might be additional transcription factors that regulate the synthesis of starch in the presence of OsSMF1 during seed development. In addition, both the ACGT and GCN4 motifs in the ONAC024 promoter provide a high-affinity binding site for OsSMF1 compared to the ACGT motif in the ONAC026 promoter. OsGZF1 is known to repress the activation of OsSMF1, in addition to promoting the down regulation of GluB-1 expression (Chen et al. 2014). Protoplast assay revealed that luciferase activity of the GZF1:Fluc vector was decreased by 0.5-fold compared with that of the negative control (Fig. 6). Expression of the RPBF transcription factor was increased in the OsSMF1-transformed calli compared with that in wild-type plants (Fig. 5), and OsSMF1 bound to the promoter of RPBF in vivo, as determined by protoplast assay (Fig. 6). Additionally, RPBF and OsSMF1 have been reported to synergistically activate transcription from the promoters of rice SSPs (Kawakatsu et al. 2009; Yamamoto et al. 2006). Knock-out transgenic rice, in which the accumulation of OsSMF1 (called RISBZ1) and RPBF was reduced, showed a significant reduction in SSPs (Kawakatsu et al. 2009). These results suggest that OsSMF1 directly regulates the expression of OsGZF1 and RPBF by binding to the ATGA motif in their promoters to regulate SSPs. However, neither OsSMF1 nor MBD1 was expressed by OsSMF1 in the protoplast assay (data not shown). The protoplast in vivo analysis revealed that the binding of OsSMF1 alone to the binding motif was not sufficient to self-activate the OsSMF1 transcription factor (data not shown). This result eliminated the possibility that OsSMF1 autoregulates itself as observed with GUS reporter genes under the control of the OsSMF1 promoter in rice protoplast (Onodera et al. 2001).

The majority of the newly identified target genes have unknown functions in seed development. We showed that two unknown target genes were highly expressed in OsSMF1-transformed calli compared with wild-type calli. Among them, OsEnS44 was identified as an OsSMF1 target gene. Interestingly, we determined that OsSMF1 bound to the promoter of OsEnS44 by protoplast transactivation assay (Fig. 6). OsEnS44 contains a thioredoxin domain, which acts in redox regulation throughout the life cycle of the seed (Wong et al. 2003). The activation of OsEnS44 by OsSMF1 may be one transcription factor cascade or may act as a transcription factor for the activation of downstream genes that regulate the biosynthesis of the major seed components. Therefore, these results provide novel information regarding the regulatory roles of OsSMF1 in seed maturation. It can be concluded that OsSMF1 is one of the key transcription factors involved in grain filling that functions by regulating the expression of a wide variety of genes during seed maturation.

**Conclusions**

Overall, the Q9-PBM results showed that OsSMF1 bound to TGAGTCA (GCN4 motif), CCACGTC (ACGT motif), and GGATGAC (ATGA motif) with different affinities. First, the transcription factor OsSMF1 regulated the expression of RPBF and OsGZF1, which contained the ATGA motif on their promoters. This result suggested that OsSMF1 regulated SSPs by binding the promoter of these genes. Second, OsSMF1 also regulated starch synthesis by binding to the promoters of ONAC024 and ONAC026, which contained either the
ACGT or GCN4 motifs. Finally, we identified OsEnS44 as an OsSMF1 target gene, which contained the GCN4 motif in its promoter region, but its function is unknown in seed development. Further studies on the regulatory mechanisms of OsSMF1, including phenotype observations in transgenic rice, will help advance the understanding of rice seed development.

Methods

Plasmid construction
OsSMF1 was isolated from panicles of Oryza sativa cv. Ilmi by RT-PCR prior to the heading stage using the primers indicated in Additional file 6: Table S1. The full-length coding region of the gene was cloned into a pET-DsRed expression vector (Kim et al. 2012) to generate an OsSMF1-DsRed fusion gene, thereby producing the plasmid pET-SMFRed. This vector was used for the expression and purification of the OsSMF1 protein in Escherichia coli.

The GFP sequence of Wsi18:GFP (Yi et al. 2011) was replaced with a Gateway ® cassette containing attR recombination sites flanking the ccdB gene and a chloramphenicol-resistance gene, thereby yielding the plasmid pSB-WsiGW. The OsSMF1-DsRed gene was then inserted into the pSB-WsiGW binary vector by a BP reaction following the manufacturer’s instructions (Invitrogen), generating the plasmid Wsi:SMF1Red. Finally, the constructs were introduced into Agrobacterium tumefaciens (Invitrogen), generating the plasmid Wsi:SMF1Red.

Protein expression and purification
The proteins were expressed in the E. coli strain BL21-CodonPlus (Stratagene). Cells were cultured overnight and inoculated into fresh liquid LB medium, grown at 37 °C to an OD260 of 0.6, and induced with 1 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) at 25 °C for 5 h. Cell pellets were obtained by centrifugation at 4 °C for 5 min at 5,000 g, followed by washing with cold PBS buffer containing a protease inhibitor (Roche). Then, the cells were resuspended in 5 ml lysis buffer (50 mM NaH2PO4, 300 mM NaCl, and 10 mM imidazole (pH 8.0)), sonicated five times with 15 s bursts and 45 s of cooling on ice, and centrifuged at 14,000 g for 30 min at 4 °C. The resulting supernatants were incubated with 2 ml Ni-NTA agarose (Qiagen) for 30 min, loaded into empty columns, and washed twice with 5 ml lysis buffer containing 10 mM imidazole according to the manufacturer’s instructions. Purified proteins were eluted with 0.5 ml lysis buffer containing 240 mM imidazole. Five micrograms of purified proteins, as measured using the Bradford assay, were used in each experiment.

Protein-binding microarray design
A microarray was designed as described previously and manufactured by Agilent Technology (Santa Clara, CA, USA) (Kim et al. 2009). This microarray (Q9-PBM) consisted of 232,145 quadrupled probes, including 131,072 probes for all possible 9-mers, 101,073 of which were replicated. Each 9-mer was concatenated four times, followed by a sequence complementary to a primer (5′-CGGAGTCACCTAGTCGAG-3′) and a 5-nt thymidine linker for attachment to the slide. Each microarray slide contained a total of 243,504 spots arranged in 267 columns and 912 rows. In addition to the quadruple probes, 1,474 random sequences from the yeast genome, 8,081 blank probes and 1,804 probes provided by the manufacturer were included.

Protein-binding microarray experiments
Complementary DNA was synthesized and analyzed to verify successful synthesis according to previously described methods (Kim et al. 2009). A double-stranded microarray was washed with PBS–0.01% (v/v) Triton X-100 and blocked with PBS-2% (wt/v) BSA (Sigma) for 1 h. Then, it was washed with PBS–0.1% (v/v) Tween-20, PBS–0.01% (v/v) Triton X-100 and PBS for 1 min in each solution. Next, a protein binding mixture was prepared containing 200 nM TF in PBS-2% (wt/v) BSA, 51.3 ng/μl salmon tested DNA (Sigma) and 50 μM zinc acetate. The mixture was incubated with the microarray for stabilization of and hybridization with the probes at 25 °C for 1 h. Subsequently, the microarray was washed with PBS-50 μM zinc acetate-0.5% (v/v) Tween-20, PBS-50 μM zinc acetate-0.01% Triton X-100 and PBS-50 μM zinc acetate for 2 min. Finally, fluorescence images were captured using a microarray scanner (Axon).

Motif extraction
The consensus binding sequence was determined based on the fluorescence signal strength according to previously described methods (Jung et al. 2012; Kim et al. 2009). Two independent linear models (y = ax + b) were applied to the steep left and extended right tail regions of the rank-ordered fluorescence signal distribution curve for the bound protein using R statistical language. Spots that exhibited strong fluorescence and high enrichment were subjected to alignment. These groups of sequences were visualized with SEQLOGO ‘Visualize information content of patterns’ [http://www.bioconductor.org/packages/release/bioc/html/seqLogo.html], which yielded an intensity profile figure, sequence logos and related statistical data. P-values and position weight matrices were calculated using the Wilcoxon-Mann–Whitney test.
Electrophoretic mobility shift assay (EMSA)

Biotin end-labeled and unlabeled oligonucleotides were annealed with each complimentary sequence (Additional file 6: Table S1). Five micrograms of OsSMF1 protein were incubated with 40 fmol biotin-labeled double-stranded oligonucleotides, 1 μg poly dl-dC, 1X binding buffer, 2.5% (v/v) glycerol and 0.05% (wt/v) NP-40 in a 20-μl reaction volume for 1 h at room temperature, according to the manufacturer’s instructions (Pierce). The reaction mixture was then analyzed by electrophoresis on a non-denaturing 6% acrylamide gel with 0.5X TBE buffer. Subsequently, the DNA-protein complexes in the gel were transferred to a positively charged nylon membrane by electrophoretic transfer in 0.5X TBE buffer. The reactions were conducted using various DNA substrate concentrations (0, 0.1, 0.2, 0.4, 0.8, 1.2, 2, and 6 μM) and 0.2 μg/μl OsSMF1. Binding was performed in the presence of 10 mM Tris, 50 mM KCl, 1 mM DTT (pH 7.5), 50 μg/μl poly(dl–dC), 2.5% glycerol, 0.05% NP-40, and 5 mM MgCl₂, and incubation for 1 h at room temperature. Following incubation, 2 μl of 10X EMSA loading dye was added to the reaction mixtures and loaded in an 8% polyacrylamide gel. The gel was stained with a SYBR Gold solution for 30 min and observed under UV light. DNA-binding band intensity was assessed using Gel-Pro Analyzer program, and the intensity value was used to calculate the $K_d$ value using Prism software. The data were fit to the following equation:

$$Y = \frac{B_{max} \times X}{(K_d + X)}$$

where $B_{max}$ is the maximal binding, and $K_d$ is the concentration of ligand required to reach half-maximal binding.

Agrobacterium-mediated transformation of rice

The Agrobacterium strain LBA4404 harboring Wsi:SMF1 was introduced into embryogenic rice calli (Oryza sativa L. Japonica cv. Ilmi) by Agrobacterium-mediated transformation (Sohn et al. 2006). Callus induction, co-cultivation with A. tumefaciens, and the selection of transformed calli were performed as previously described (Sohn et al. 2006). Selected calli were subcultured for 2 weeks on fresh 2 N6 medium with 250 mg/L cefotaxime, and then transferred to MSR medium containing 250 mg/L cefotaxime and 4 mg/L phosphinothricin and incubated for 4 weeks at 27 °C under continuous light for selection and regeneration. Regenerated shoots were transferred to MSO medium containing 4 mg/L phosphinothricin and incubated for 4 weeks for root induction. The plantlets were then transplanted to a Wagner pot (200 cm²) kept in a greenhouse for subsequent growth. Wsi:SMF1 transgenic rice of the T₁ generation were used in further analyses.

RNA isolation and RT-PCR

Total RNA was extracted from leaves of transgenic and wild-type rice plants using TRI REAGENT™ (Molecular Research Center, www.mrcgene.com) and purified with a Qiagen RNeasy Mini Kit (Qiagen, www.qiagen.com). cDNA templates were synthesized using RevertAid H minus M-MulV reverse transcriptase (Fermentas, www.fermentas.com). Semiquantitative RT-PCR analysis was performed was performed in a 20 μl reaction mixture under the following conditions: one cycle at 95 °C for 2 min and 25 to 35 cycles at 94 °C for 30 s, 60 °C for 30 s, and 72 °C for 30 s. Real-time quantitative RT-PCR analysis was performed using 2X Real-Time PCR Premix with EvaGreen (Solgent, www.solgent.com) according to the manufacturer’s protocol. Thermal cycling and fluorescent detection were performed using a Stratagene Mx3000P Real-Time PCR machine and Mx3000P software v2.02 (Stratagene, http://www.genomics.agilent.com). Melting curve analysis (with an increase from 55 to 65 °C in increments of 0.1 °C s⁻¹) was performed to ensure that only the desired PCR product was measured at a specific melting temperature. Real-time PCR was performed in triplicate for each cDNA sample. Following amplification, the PCR data were assessed using a comparative quantification (calibrator) analysis with Mx3000P software v2.02. (Stratagene). The rice tubulin gene (Os11g0247300) was used as an endogenous control. All primer pairs used are listed in Table S2.

Transcriptional activity assay in rice protoplast

Transcriptional activity assay of OsSMF1 in rice protoplast was conducted using a dual luciferase reporter assay system (Promega, USA). To construct reporter plasmids, promoters of putative target genes of OsSMF1 were amplified from Oryza sativa cv. Ilmi genomic DNA using specific primers (Additional file 6: Table S1) and were then cloned into a pHBT vector (GenBank accession no. EF090408) between the firefly luciferase (Fluc) gene and nos terminator, respectively. For dual luciferase assay, the OsSMF1 and Renilla luciferase (rLUC) coding sequences were cloned into the pHBT vector between the 35S promoter and nos terminator, respectively.

The resulting construct, P35S:OsSMF1, was co-transformed with the reporter plasmid into isolated rice protoplasts by polyethylene glycol (PEG)-mediated transformation (1 μg per transfection). P35S:rLUC (Renilla luciferase) was also added to each sample as an
endogenous control (1 μg per transfection). Protoplast isolation and PEG-mediated transformation were performed as previously described (Jung et al. 2015). Luciferase activity was detected using an Infinite® 200 (Tecan, Switzerland). Measured fLUC activities were normalized to rLUC activities.

Statistical analyses
All statistical analyses were performed using Sigma Plot v10 software.

Additional files

Additional file 1: Figure S1. Transcript levels of Wsi18, OsSMF1, and OsRDM from 300 K Rice Genome Microarray (www.ggbio.com). The transcript levels were measured in different sized panicles before heading (1, 3, 5, 8, 10, 15, 20, and 22 cm), at the indicated days after pollination (1, 3, 4, 11, and 21 days) and in the leaf, root, germinating seed, callus, and regenerating callus. (PPTX 70 kb)

Additional file 2: Figure S2. Rank analysis of OsSMF1 binding by Q9-PBMW analysis. According to the rank-ordered signal distribution, two independent linear models (y = ax + b) were applied in the deep (b1 = 5032.03, slope = −38.3) and heavy right (b1 = 978.4, slope = −0.00657) tall regions of the curve. The extrapolated rank estimation for motif extraction was 1.286. (PPTX 43 kb)

Additional file 3: Table S2. Primers used for PCR/real-time PCR. (XLSX 12 kb)

Additional file 4: Figure S3. The query gene, OsSMF1, is marked by an asterisk. Each circle indicates a gene, and the lines represent the correlations between the genes. Eighty-five genes were identified as OsSMF1-related genes, with a minimum correlation value of 0.55 and depth of 1. (PPTX 70 kb)

Additional file 5: Figure S4. The expression patterns of transcription factors as a putative target of OsSMF1 based on the 300 K Rice Genome Microarray (www.ggbio.com). The expression was measured in different sized panicles before heading (1, 3, 5, 8, 10, 15, 20, and 22 cm), at the indicated days after pollination (1, 3, 4, 11, and 21 days) and in the leaf, root, germinating seed, callus, and regenerating callus. (PPTX 75 kb)

Additional file 6: Table S1. List of putative OsSMF1 target genes, as determined by RAN analysis (r-value ≥ 0.55, depth = 1). (XLSX 19 kb)

Abbreviations
OsSMF1: A basic leucine zipper transcription factor involved in the regulation of rice seed maturation; PBM: Protein-binding microarrays; RAN: RiceArrayNet; SSF: Seed storage protein

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