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(71) Applicant (for all designated States except US): **NATIONAL RESEARCH COUNCIL OF CANADA** [CA/CA]; 1200 Montreal Road, Ottawa, Ontario K1A 0R6 (CA).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **SELVARAJ, Gopalan** [CA/CA]; 540 Nesslin Crescent, Saskatoon, Saskatchewan S7J 4V5 (CA). **WANG, Aiming** [CA/CA]; 571 Mapledale Ave., London, Ontario N5X 1H3 (CA). **XIA, Qun** [CA/CA]; 2119 Kenderdine Road, Saskatoon, Saskatchewan S7N 4A5 (CA). **XIE, Wenshuang** [CA/US];

5529 Colusa Avenue, Richmond, CA 94804-5227 (US). **DATLA, Raju** [CA/CA]; 527 Bayview Terrace, Saskatoon, Saskatchewan S7V 1B6 (CA).

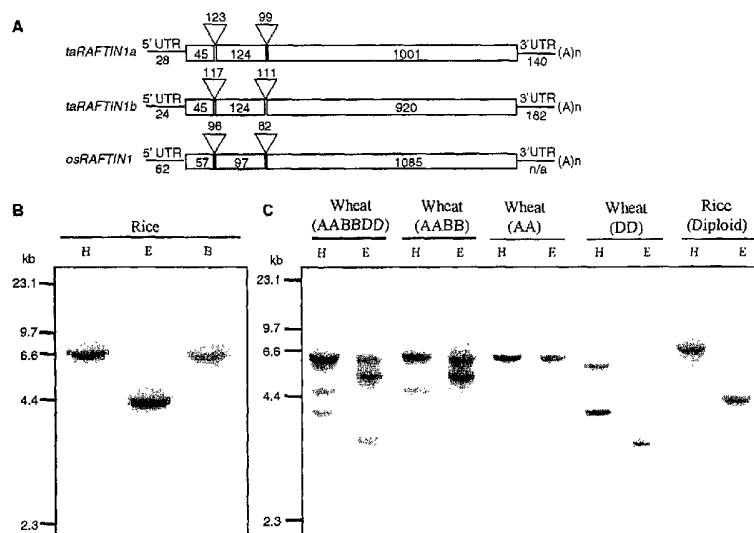
(74) Agents: **CONN, David, L.** et al.; Borden Ladner Gervais LLP, World Exchange Plaza, 100 Queen Street, Suite 1100, Ottawa, Ontario K1P 1J9 (CA).

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(54) Title: **RAFTIN GENE, PRODUCT, AND USE THEREOF**



(57) Abstract: The anther recruits a number of specific genes to support microspore development. Here we report identification of a novel gene, RAFTIN1, from bread wheat and rice, encoding a protein containing a BURP domain hallmarked with 4 CH repeats at its C-terminus. This single copy gene (per haploid complement in cereals) is exclusively expressed in the tapetum during post-meiotic stages when the young microspore undergoes rapid expansion. RAFTIN1 is biosynthesized in the tapetum, transported into the Ubisch body, and further deposited onto the microspore exinewall. Transgenic rice, in which RAFTIN1 expression is down-regulated, shows normal growth and development but also shows male sterility. In the RAFTIN1-less anther, tapetal degeneration is retarded and pollen lacks contents. Thus, RAFTIN1 assembly in the Ubisch body and the microspore exinewall is required for microspore development, probably for regulation of metabolite transport from the tapetum to the microspore. Silencing or knocking-out of RAFTIN will find utility in breeding programs wherever male sterile lines are required.



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***RAFTIN* GENE, PRODUCT, AND USE THEREOF**

TECHNICAL FIELD

5 This invention generally relates to a method of modulating male fertility in plants, and more specifically relates to a method of modulating male fertility in cereal plants by down regulating expression of *RAFTIN* genes. Modulation of male fertility in plants is of increasing significance and utility in controlled plant breeding programs in which male sterile lines are useful. Male sterile lines are of interest and use commercially and can be
10 useful in reducing unwanted gene flow among different breeds of the same species of plant, for example from a transgenic crop to its non-transgenic neighbours.

BACKGROUND

 The anther, a tiny, short-lived but functionally complex organ encompassing the
15 male gametophyte plays a fundamental role in the reproductive cycle of flowering plants as well as in the agricultural practice, i.e. hybrid seed production. As one of the most elaborate and complex processes in plant life, anther development, in particular at the molecular level still remains poorly understood. Previously cytological observations through the light and electron microscopes have sketched a generalized scheme of
20 developmental events leading to the ontogeny of the anther architecture for most angiosperms (D'Arcy, 1996, Goldberg et al., 1993, Shivanna et al., 1997). Differentiating from stamen primordia, the anther initially consists of a mass of homogeneous cells surrounded by the epidermis. Followed with the differentiation of archesporial cells in the hypodermal region, a four-lobed structure (four microsporangia or locules) is developed.
25 A periclinal division of archesporial cells gives rise to the formation of the outer, primary parietal layer and the inner, sporogenous layer. The former, upon cell divisions generates multi-layer cells which are differentiated into the endothecium (the outermost layer beneath the epidermis), the middle layer (one to three layers between the endothelium and the tapetum) and the tapetum (the innermost layer). Concomitant with the differentiation
30 of the anther wall (the epidermis, the endothecium and the middle layer) and the tapetum, the centered sporogenous layer gives rise to microspore mother cells (MMCs). The differentiated MMC undergoes meiosis to produce tetrads of microspores. Further development from young microspores to mature pollen grains, morphologically evident

with the enlargement of the anther and microspores, the degeneration of the tapetum, pollen desiccation, anther dehiscence and pollen release, is an exquisite interplay between the male gametophyte and sporophytic tissues of the anther (Goldberg et al., 1993, McCormick, 1993).

5 A rapidly increasing body of evidence suggests that the development of normal pollen grains requires a functional tapetum (Aarts et al., 1997, De Block et al., 1997, Goldberg et al., 1993, Mariani et al., 1992, Mizelle et al., 1989). As early as prior to the onset of meiosis, the tapetum tissue initially networks meitocytes via plasmodesmata with the rest of the supporting tissues of the anther. Upon the formation of postmeiotic tetrads,
10 the tapetum functions as nursing cells to produce callase to disassociate the tetrads. During the following free microspore and vacuolated microspore stages, the tapetum supplies essential nutrients and metabolites to assist in microspore expansion, vacuolation and extracellular matrix formation (Mascarenhas, 1990, Mizelle et al., 1989, Sanders et al., 1999). Towards the end of the vacuolated microspore stage, the tapetum triggers
15 programmed degeneration that continues throughout the vacuolated pollen grain stage, releasing its macromolecules such as proteins, lipids and carbohydrates, some of them utilized in the build-up of the complex extracellular wall surrounding the microspore (Furness & Rudall, 2001, Mascarenhas, 1975, McCormick, 1993, Mizelle et al., 1989, Piffanelli et al., 1997, Shivanna et al., 1997, Wiermann & Gubatz, 1992). However, by
20 what means the tapetal metabolites are transported onto the surface of microspore is obscure and how the sporophytically produced, gametophytically localized proteins contribute to microspore development is not known.

In this study, we report the isolation of three apparently homologous anther-specific genes *taRAFTIN1a*, *taRAFTIN1b* and *taRAFTIN1d* in allohexaploid wheat
25 (*Triticum aestivum* L.) and their ortholog *osRAFTIN1* in rice (*Oryza sativa* L.). We provide evidence that *RAFTIN1* transcription only took place in the tapetum but their proteins were predominantly evident in the tapetum, the Ubisch body and the microspore exinewall, establishing an example of the tapetum-Ubisch Body-microspore transport pathway. Our data show that *RAFTIN1*, whose structural counterpart is not found in the
30 *Arabidopsis* genome or other eudicot genomes but found in the ESTs derived from the anther or anther-containing reproductive tissues of various monocot species including barley, sorghum, hexaploid wheat, rice, wild diploid wheats, maize and rye contains a BURP domain that has been only found in the plant kingdom, thus constituting the first

anther-specific version of BURP domain proteins. Our results demonstrate that in the transgenic rice where *RAFTIN1* expression was down-regulated, the normal tapetal degeneration was retarded and microspore contents were lacking, leading to the production of male sterile pollen. We suggest that the Ubisch body- and microspore exinewall-assembled *RAFTIN1* proteins are probably involved in the transport of the metabolites from the tapetum to microspore, that is required for anther development in rice, wheat and probably in other cereal species. This will prove useful in manipulating male sterility in such other plants such as species of monocotyledon, e.g. rye, oats, barley, sorghum and maize (in addition to wheat and rice).

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1. Genomic Organization of *RAFTIN1* Gene Family in Wheat and Rice.

(A) Schematic representation of genomic structures of cloned *RAFTIN1* genes in wheat and rice. *taRAFTIN1a* and *taRAFTIN1b* were isolated from a hexaploid wheat, and *osRAFTIN1* from rice. The numbers denote nt sequence length of the untranslated regions (UTR), exons (boxes) and introns (triangles). Not drawn to scale. (B) *RAFTIN1* gene in rice. Southern blot of rice genomic DNA (~10 µg) digested with *HindIII* (H), *EcoRI* (E) and *BamHI* (B) and probed with the entire *osRAFTIN1* ORF retrieved by PCR; size markers are from a λ DNA-*HindIII* digest. (C) *taRAFTIN1a* and closely related sequences in hexaploid, tetraploid and diploid wheats, and diploid rice. Southern blot of genomic DNA (~15 µg for wheat and ~10 µg for rice) restricted with *HindIII* (H) and *EcoRI* (E) and probed with the entire *taRAFTIN1a* ORF retrieved by PCR; size markers are from a λ DNA-*HindIII* digest.

Figure 2. Anther-Specific Expression of *RAFTIN1* in Wheat and Rice. (A) and (C) Northern blot analysis of *RAFTIN1* expression in hexaploid wheat and rice. Total RNA (5 µg) from each tissue was loaded. (A) probed with the entire *taRAFTIN1a* ORF retrieved by PCR. (C) probed with the entire *osRAFTIN1* ORF retrieved by PCR. (B) and (D) RT-PCR analysis of *RAFTIN1* expression in wheat and rice. (B) in wheat. (D) in rice. Primers and PCR conditions were described in METHODS. Total RNA isolated from different tissues indicated here was used for RT-PCR and northern blotting analyses. Root, root tissue; Stem, stem tissue; Leaf, leaf tissue; Flower, developing young flower tissue; Fl w/o anther, developing young flower tissue with anther removed; Anther, developing anther tissue. For northern blot analyses, the panel underneath is the same gel used for

blotting stained with ethidium bromide. For RT-PCR, the panel underneath is the control RT-PCR of a housekeeping gene, GAPDH (glyceradehyde-3-phosphate dehydrogenase, taGAPDH for wheat, osGAPDH for rice).

Figure 3. Detection of RAFTIN1 Proteins in Wheat Anther. Protein extraction and western blot analysis were performed as described in METHODS. Purified taRAFTIN1a antiserum IgG was used for detection. Total proteins were extracted from different wheat tissues as indicated in Figure 3.

Figure 4. Amino Acid Sequences of RAFTIN1 Proteins and BURP Domains. (A) Comparison of the deduced primary structure of *RAFTIN1* gene products. The predicted transmembrane domains are indicated as a filled bar. The specific sequences of these regions in each gene product are boxed. (B) Alignment of BURP domains. Substitutions are shown as such: identity as hyphens, and the gaps introduced in the alignment as triangles. The highly conserved amino acids, and CH motifs are indicated as dots and stars, respectively.

Figure 5. *In situ* RNA Hybridization of *RAFTIN1* Transcripts and Immunocytochemical Detection of RAFTIN1 Proteins in Cross-Sections of Hexaploid Wheat Flowers. From (A) to (D), detection of *RAFTIN1* mRNA in wheat young florescence. (A) Hybridized to a *taRAFTIN1a* antisense probe. (B) Hybridized to a *taRAFTIN1a* sense probe (control). (C) and (D) Enlargement of (A) and (B), respectively. (E) and (F) Detection of *RAFTIN1* mRNA in rice young florescence. (E) Hybridized to an *osRAFTIN1* antisense probe. (F) Hybridized to an *osRAFTIN1* sense probe (control). From (G) to (M), detection of RAFTIN1 proteins in wheat young inflorescence by taRAFTIN1a polyclonal antibodies, except for (H) which has been probed with pre-immune sera (control). From (I) to (J), the sections showing various developmental stages, progressing from the left to the right, were chosen. (I), "tetrad" stage; (J), "free microspore" stage; (K), "vacuolated microspore" stage; (L), "vacuolated pollen grain" stage; (M), "3-nucleate pollen grain" stage. Purified taRAFTIN1a antiserum IgG was used for detection. The bluish precipitate due to positive reaction is indicated by solid arrows. tp: tapetum; ms, microspore; ps; pollen sac; pg: pollen grain; ep: epidermis; en: endothecium. In (A) and (B), scale bars=200 μ m; in (C) and (D), =100 μ m; all other panels: 40 μ m.

Figure 6. Electron Micrographs of Wheat Anther Tissues Labelled with Polyclonal Antibodies Raised against taRAFTIN1a. (A) Anther ultra-thin sections immunolabeled with anti-taRAFTIN1a IgG. x7531. (B) Tapetal cells and Ubisch bodies

immunolabeled with anti-taRAFTIN1a IgG. x45500. (C) Exinewall of microspore immunolabeled with anti-taRAFTIN1a IgG. x29640. ep, epidermis; en, endothecium; ml, middle layer; ta, tapetum; ub, Ubisch body; ex, exine; ms, microspore; ob, orbicular wall; ba, bacule; te, tectum; and fl, foot layer.

Figure 7. Silencing *osRAFTIN1* in Transgenic Rice using an Intron-Containing Hairpin RNA Strategy. (A) Schematic representation of intron-containing hairpin constructs designed for genetic transformation. (B) Northern blot analysis of *osRAFTIN1* expression in transgenic rice transformed with intron-containing hairpin constructs. Percentage of *osRAFTIN1* expression in different lines relative to that of WT normalized against total RNA loaded is given.

Figure 8. Phenotypes of Wild-Type Rice and *osRAFTIN1*-Silenced Lines. (A) Vegetative growth of a wild-type rice line (457-5) transformed with a hygromycin resistant gene. (B) Vegetative growth of a transgenic line (507-5) transformed with a *35S::hy-osRAFTIN1* chimeric gene. (C) Mature florescence of Line 457-5. (D) Mature florescence of Line 507-5. (E) Rice grains of Line 457-5. (F) Rice grains (empty) of Line 507-5. (G) Hulls and rice of (E). (H) Hulls and a few undeveloped embryos of (F). (I) Rice grains of Line 507-5 fertilized with wild-type pollen.

Figure 9. Scanning Electron Micrographs of Anthers and Pollen Grains of Wild-Type Rice and *osRAFTIN1*-Silenced Lines. (A) A representative of mature anthers from wild-type rice. (B) A representative of pollen grains from a mature undehisced wild-type anther. (C) Surface of a mature pollen grain from a wild-type anther. (D) A representative of mature pollens from transgenic Line 507-5 in which *osRAFTIN1* anther-specific expression was down-regulated by an intron-spliced hairpin RNA strategy. (E) A representative of pollen grains from a mature undehisced anther in Line 507-5. (F) Surface of a mature pollen grain from Line 507-5.

In (A) and (D), scale bars=1 mm; in (B) and (E), =10 μ m; in (C) and (F), =1 μ m.

Figure 10. Transmission Electron Micrographs of Anthers and Pollen Grains of Wild-Type Rice and *osRAFTIN1*-Silenced Lines. (A) and (B) Wild-type. (C) and (D) *osRAFTIN1*-silenced line 507-5. Scale bars=10 μ m. ep, epidermis; en, endothecium; ta, tapetum; ub, Ubisch body; ex, exine; ms, microspore; ob, orbicular wall.

Figure 11. *taRAFTIN1a* cDNA sequence (1338 nt excluding the polyA tail, ORF from nt 29 to nt 1198) (SEQ ID NO: 29). Start codon and stop codon are underlined.

Figure 12. *taRAFTIN1a* genomic sequence (1560 bps including two introns). Introns are shown in lower case letters (SEQ ID NO: 30). Start codon and stop codon are underlined.

Figure 13. *taRAFTIN1a* promoter sequence (1719 bps) (SEQ ID NO: 31).

5 **Figure 14.** *taRAFTIN1b* cDNA sequence (1275 bps excluding the polyA tail, ORF from nt 25 to nt 1113) (SEQ ID NO: 32). Start codon and stop codon are underlined.

Figure 15. *taRAFTIN1b* genomic sequence (1503 bps including two introns). Introns are shown in lower case letters (SEQ ID NO: 33). Start codon and stop codon are underlined.

10 **Figure 16.** *taRAFTIN1b* promoter sequence (2095 bps) (SEQ ID NO: 34).

Figure 17. *taRAFTIN1d* predicted cDNA sequence (246 bps) (SEQ ID NO: 35).

Figure 18. *taRAFTIN1d* partial genomic sequence (441 bps) (SEQ ID NO: 36).

Introns are shown in lower case letters.

Figure 19. *osRAFTIN1* cDNA sequence (1301 bps, ORF from nt 63 to nt1301) (SEQ ID NO: 37). Start codon and stop codon are underlined.

Figure 20. *osRAFTIN1* genomic sequence (1479 bps, two introns included) (SEQ ID NO: 38). Introns are shown in lower case letters.

Figure 21. *osRAFTIN1* promoter sequence (1461 bps) (SEQ ID NO: 39).

Figure 22 represents predicted protein sequences of *taRAFTIN1a* (389 residues) (SEQ ID NO: 40), *taRAFTIN1b* (362 residues) (SEQ ID NO: 41), *taRAFTIN1d* (partial sequence, 82 residues) (SEQ ID NO: 42) and *osRAFTIN1* (412 residues) (SEQ ID NO: 43).

DETAILED DESCRIPTION OF THE INVENTION

25 Genomic Organization of *RAFTIN1* in Hexaploid Wheat and Diploid Rice

An anther-specific cDNA clone (A71) encoding a polypeptide was obtained from a wheat anther full-length cDNA library. Further 3 screens of the library using the 5' region of the A71 as a probe identified, based on their sequence differences in 5' and 3' regions, two groups of cDNAs designated as *taRAFTIN1a* and *taRAFTIN1b*, respectively. The longest from each group, found to be full-length cDNA clones by the 5' rapid amplification of cDNA ends (RACE) analysis, were completely sequenced. *TaRAFTIN1a* was 1356 nucleotides (nt) in length, and had a predicted open reading frame of 1170 nt (from nt 29 to 1198) encoding a protein with 389 amino acids (aa) (41.3 kDa) while

taRAFTIN1b had a shorter cDNA (1285 nt) and a smaller ORF (1089 nt, from nt 25 to 1113) coding for 362 aa (38.3 kDa). Pairwise BLAST analysis of these two ORFs showed that they were 93% identical to each other at the nt level and 83% at the aa level. The corresponding genomic sequences of *taRAFTIN1a* and *taRAFTIN1b* cDNAs were
5 retrieved from the wheat genome using a polymerase chain reaction (PCR) approach. Comparison of *RAFTIN1* cDNA sequences with their corresponding genomic sequences revealed that both transcripts of *taRAFTIN1a* and *taRAFTIN1b* were interrupted by two introns in the 5' regions of the ORFs and composed of 3 exons (**Figure 1A**). During an endeavor to obtain more *taRAFTIN1a* and *taRAFTIN1b* related genes, a 701-bp genomic
10 fragment (named *taRAFTIN1d*) whose predicted cDNA (246 bp) was 96% and 97% identical to the 5' terminal coding regions of *taRAFTIN1a* and *taRAFTIN1b*, respectively, was isolated from the hexaploid wheat by PCR, whereas three additional screens of the anther cDNA library failed to isolate *taRAFTIN1d* cDNA or identify any new *RAFTIN1-like* genes. Since *taRAFTIN1d* had only the partial genomic sequence, it was not
15 characterized further in this study.

As BLASTN searches of the public databases only identified a hypothetical gene (AP000364) obtained from the rice genome sequencing, hereafter named *osRAFTIN1*, the predicted coding region of *osRAFTIN1* cDNA and its corresponding genomic DNA were
20 cloned by RT-PCR and PCR, respectively. The ORF of *osRAFTIN1*, also interrupted by two introns at its 5' terminal region is 1239 nt in length, encoding a 412 aa polypeptide with a predicted molecular mass of 42.3 kDa *osRAFTIN* ORF cDNA shared ~ 66% identity to *taRAFTIN1a* and *taRAFTIN1b* ORFs. The polypeptide of *osRAFTIN1* was overall ~58% identical to *taRAFTIN1a* and *taRAFTIN1b*, and had a similar size and structure.

25 DNA gel blot analysis was performed to estimate the copy number of *RAFTIN1* in rice and in the allohexaploid (AABBDD genome), tetraploid (AABB) and diploid wheats (AA or DD) using the coding region of *osRAFTIN1* or *taRAFTIN1* as a probe. The *osRAFTIN1* probe hybridized to only one DNA fragment of the rice genome digested with each of 3 different restriction enzymes that do not cleave the coding region (**Figure 1B**).
30 All recognized bands were of the sizes consistent with those obtained through electronic southern analysis of the genomic sequences retrieved from the public domain (GenBank accession no: AP000364, Sasaki, T., Matsumoto, T. and Yamamoto, K., 1999). Thus, there only exists one copy of *RAFTIN1* gene in the rice genome. When the same digestion

was probed with the wheat *taRAFTIN1a* ORF cDNA, the probe specifically hybridized to the DNA fragments with the same sizes as recognized by the *osRAFTIN1* probe (Figures 1B and 1C), confirming a close relationship between rice *osRAFTIN1* and wheat *RAFTIN1* genes. In wheat as shown in Figure 1C, there were 3 to 4 bands in the hexaploid (AABBDD genome), 2 in the tetraploid (AABB), 1 in one diploid (AA), and 1 to 2 in another diploid (DD). Careful comparison of the size of band(s) among different genomes strongly suggested that the *RAFTIN1* gene family was most likely represented by one copy per A or B or D complement in wheats, the isolated *taRAFTIN1a* and *taRAFTIN1b* genes were likely from the AABB genome and the related gene in DD genome might consist of one *HindIII* restriction site in the coding region resulting two hybridization bands. This assumption was partially supported by the observation that a closely related genomic clone isolated from an AA genome wheat was closely related to *taRAFTIN1a* and the EST clone (GenBank accession no: BG274249, Anderson, O., 2001) obtained from a BB genome wheat was to *taRAFTIN1b* (data not shown).

***RAFTIN1* Is Specifically Expressed in the Anther**

The expression pattern of *RAFTIN1* in wheat and rice was determined by probing RNA blots with ORFs of *taRAFTIN1a* and *osRAFTIN1*. The *taRAFTIN1a* probe strongly hybridized only to RNA isolated from wheat young florescence or the anther but not from root, stem, leaf or the young florescence whose anther had been removed (**Figure 2A**).

Hybridization of transcripts isolated from corresponding tissues in rice using the *osRAFTIN1* probe produced comparable results (**Figure 2C**). In both species, the single hybridization band of ~1.3 kb was consistent with the expected sizes of the *RAFTIN1* cDNAs (**Figures 2A and 2C**). For further confirmation, more sensitive RT-PCR analysis was carried out with primer pairs designed to cover 2 partial exons containing 1 intron in-between in the genomic (for wheat *RAFTIN1a*) or containing 1 entire exon flanked with two intron (for rice *osRAFTIN1*) in-between in the genomic so as to discriminate amplicons of mRNA (875 bp for wheat; 441 bp for rice) and genomic DNA origin (974 bp for wheat; 703 bp for rice). A strong band with the predicted sizes of amplified cDNA fragments (875 bp for wheat; 441 bp for rice) was detected in the young florescence or anther samples starting from 22 cycles under the given conditions (see METHODS) but not (even up to 35 cycles) in the samples of root, stem, leaf or the florescence whose anther had been removed (**Figures 2B and 2D**). Thus, *RAFTIN1* transcription was stringently restricted in the anther tissue.

To detect RAFTIN1 proteins *in planta*, the polyclonal antibodies were raised against the N-terminal portion of the RAFTIN1a fusion protein over-expressed in *E. coli*. The purified antibodies were used for Western blot analysis of different wheat tissues. Consistently, RAFTIN1 proteins were only evident in wheat young florescence or the
5 anther but not in root, stem, leaf or the young florescence whose anther had been removed (F1 w/o anther) (**Figure 3**). The detected protein was ~40 kDa in size, close to the predicted molecular weights (41.3 kDa and 38.3 kDa) for *taRAFTIN1a* and *taRAFTIN1b*. These results further confirmed the presence of *taRAFTIN1a* protein in the wheat anther.

***RAFTIN1* Encodes a BURP Domain Containing Protein and *RAFTIN1*-like Gene Are
10 not Present in the *Arabidopsis* Genome**

Since the *Arabidopsis* genome has been sequenced and large knockout population is available, we searched the genomic sequence for *RAFTIN1*-like genes (www.arabidopsis.org). BLAST analysis did not identify in the model plant any known or hypothetical genes statistically significantly homologous to *RAFTIN1* nt sequences. No
15 overall significantly similar proteins were found in the *Arabidopsis* genome by BLASTP analyses. However, limited homology was shown between the C-terminal moiety (~200 a.a.) of *RAFTIN1* and the C-termini of 5 putative gene products, 36% identical to RD22 (a.a. 175-389) (Yamaguchi-Shinozaki & Shinozaki, 1993), 35% to an unknown protein (a.a. 86-277) (F13F21.25), 30% to an aromatic rich glycoprotein (a.a. 411-588) (F1707.9),
20 26% to a putative polygalacturonase isoenzyme 1 beta subunit (a.a. 311-619) (F508.31) and 27% to another polygalacturonase isoenzyme 1 beta subunit (a.a. 390-620) (T13D8.26). These conserved homologous regions were previously named BURP domain (see below). Searches for ESTs of these genes did not reveal any evidence that any EST of them was derived from anther or florescence tissues. Thus, apparently the rice and
25 wheat anther-specific *RAFTIN1* homologues were not present in the *Arabidopsis* genome.

Further BLASTX searches of the public gene databases for *RAFTIN1* homologous sequences corroborated that there were no overall similar genes documented in plants other than monocot cereals. In cereals, the significant matches were shown to 12 ESTs (~600 bp) from anthers, young panicles or pre-anthesis spikes in bread wheat, a wild
30 diploid wheat (*Aegilopes speltoides*, BB genome), rice, barley, rye, sorghum and maize. The GenBank accession numbers of these ESTs containing the longest 5' sequences of each identical gene included BE40071 (hexaploid wheat, available sequence identical to the 5' terminal region of *taRAFTIN1a* cDNA), BE499238 (hexaploid wheat, identical to

the 5' terminal region of *taRAFTIN1b*), BG274249 (*Aegilops speltoides*, similar to the N-terminus of *RAFTIN1*), AU029260 (rice, identical to *osRAFTIN1*), AW562783 (maize, similar to the N-terminus of *RAFTIN1*), BE060637 (barley, similar to the N-terminus of *RAFTIN1*), BE636918 (rye, similar to the N-terminus of *RAFTIN1*) and BI140560

(sorghum, similar to the N-terminus of *RAFTIN1*). Domain searches revealed that there were two predicted transmembrane domains localized at the very N-terminal and the central regions, and a BURP domain at their C-terminal moiety (**Figure 4A**). The BURP domain conserved in diverse plant proteins has recently been suggested to play an important and fundamental but uncharacterized role in plant (Hattori et al., 1998). The sequences of BURP domains, identified in this study and previously, were compared. Remarkably, all BURP domains conserved two phenylalanine residues (FF) at their N-termini of the domain and four repeated cysteine-histidine (CH) motifs in the consensus sequence CHX₁₀CHX₂₅₋₂₇CHX₂₄₋₂₅CH, where X represents any amino acid (**Figure 4B**), similar with the previous finding (Hattori et al., 1998).

***RAFTIN1* Transcripts Are Found only in the Tapetum but *RAFTIN1* Proteins Are Present in the Tapetum and Microspore during the Rapid Growth Stages of Microspore**

The anther encloses morphologically and functionally divergent tissues such as the anther wall (the epidermis, the endothecium and the middle layer), the tapetum, microspore and other supportive tissues. *RAFTIN1* transcripts were localized by *in situ* RNA hybridization using a stretch of *taRAFTIN1a* or *osRAFTIN1* antisense RNA synthesized *in vitro* as a probe. In both wheat and rice, the *RAFTIN1* mRNA was found only in the tapetum but not in the ovary, the anther wall, microspore, the filament and other supportive tissues (**Figures 5A, 5C and 5E**). The hybridization signal was negative when the *taRAFTIN1a* or *osRAFTIN1* sense RNA was used as a probe (**Figure 5B, 5D and 5F**). Thus, *RAFTIN1* transcription was confined to the mono-layer cells of the tapetum surrounding microspore.

Cellular localization of *RAFTIN1* proteins in the wheat anther was conducted by immunocytochemical analysis using the *taRAFTIN1a* polyclonal antibodies. Surprisingly, the positive signal for the presence of *RAFTIN1* proteins was not only evident in the tapetum but also in microspore, albeit not detected in other tissues (**Figure 5G**). The fidelity of this signal was corroborated by a pre-serum control (**Figure 5H**). The detection of *RAFTIN1* protein into the tapetum and microspore was inconsistent with tapetal

localization of *RAFTIN1* transcripts arising from the *in situ* RNA localization. This observation raised the possibility for translocation of RAFTIN1 proteins from tapetal cells to microspore.

To further explore the temporal expression pattern of *RAFTIN1*, the wheat floral sections at the various developing stages of microspore were probed with the purified taRAFTIN1a antibodies. RAFTIN1 proteins were not evident before the tetrad stage (**Figure 5I**) and strongly detected in both the tapetum and the male gametophyte during post tetrad stages from the young free microspore stage to the vacuolated pollen grain stage when microspore underwent rapid expansion (**Figures 5J and 5K**). And weak signals were also found in the pollen maturation stages (**5L and 5M**). This spatio-temporal expression pattern of *RAFTIN1* was further confirmed by histochemical GUS assay of the transgenic tobacco, *Arabidopsis* and rice plants transformed with an *RAFTIN1* (*taRAFTIN1a*, *taRAFTIN1b* or *osRAFTIN1*) promoter: *GUS* chimeric gene (data not shown). These results indicated that *RAFTIN1* transcription and translation were highly regulated during the male gametophyte development in wheat and rice.

RAFTIN1 Proteins Are Subcellularly Localized to the Ubisch Body and the Microspore Extracellular Wall

The taRAFTIN1a antibodies were used for subcellular localization of RAFTIN1 by immunoelectronic microscopy analysis. Consistent with the results obtained from the immunocytochemical assay, the antibodies exclusively detected the presence of RAFTIN1 in the tapetum and microspore (**Figure 6A**). However, no significant labeling was found in the epidermis, the endothecium and other anther supportive tissues (**Figure 6A**). No labeling was detected anywhere with the purified preimmune serum. Examinations of the positive cells immunolabeled with the anti-taRAFTIN1a serum revealed that gold labeling was most evident in the tapetal microsome-like structures, the Ubisch body, and the extracellular matrix (exinewall) of microspore but not in the orbicular wall or inside microspore (**Figures 6B and 6C**), further supporting that microspore was not likely to be the site for RAFTIN1 production. In the exinewall, gold particles were clearly shown in the tectum, the foot layer and bacules (**Figure 6C**). Although lacking direct evidence, Ubisch bodies were previously hypothesized to be involved in transport of sporopollenin from the tapetum to the developing microspores (Huysmans et al., 1998). The localization of RAFTIN1 proteins in Ubisch bodies and the physical connection between the tapetum and the Ubisch body and between the Ubisch body and microspore supported that Ubisch

bodies transported RAFTIN1, probably in a fashion similar to that by which sporopollenin is transported.

Intron-Spliced Hairpin RNA Effectively Reduces *osRAFTIN1* Expression in Rice

RAFTIN1 is a novel protein with no established function and is without a structural counterpart in *Arabidopsis*. Taking advantage of the findings that there is only one copy of *RAFTIN1* in rice (in this study) and an intron-spliced hairpin RNA (ihpRNA) can efficiently induce sequence-specific gene silencing in plants (Smith et al., 2000), the transgenic approach was employed to silence *osRAFTIN1* in rice to explore *RAFTIN1* functions *in planta*. Eight rice transformation vectors harboring DNA sequences encoding ihpRNA homologous to *osRAFTIN1* sequence under the control of *taRAFTIN1a*, *taRAFTIN1b*, *osRAFTIN1* or *35S* promoters were constructed (**Figure 7A**); the chimeric genes were transformed into rice. Fifty-four lines transformed with these constructs and nine lines transformed with a selection marker gene (as a control) were recovered, showing resistance to the selection antibiotic, hygromycin and being positive in the PCR screening (data not shown). RNA isolated from the anthers of one randomly chosen representative line of each construct was used for Northern blot analysis to evaluate silencing efficacy (**Figure 7B**). Of the transgenic lines examined, a line (497) containing *osRAFTIN1* promoter, a partial genomic portion including an intron (at 3' end) of *osRAFTIN1*, and the corresponding antisense sequence of *osRAFTIN1* achieved the best silencing efficacy, with only 3.7% intact transcripts detected, in comparison with that in a control line (WT) containing the selection marker gene (**Figure 7B**). The transgenic line with the poorest reduction of *osRAFTIN1* expression was transformed with a *35S* promoter followed with a *taRAFTIN1a* hairpin DNA, silencing 37.6% intact transcripts relative to WT. Thus, all the designed chimeric genes in the randomly picked lines could effectively induce *osRAFTIN1* gene-specific silencing in the transgenic rice.

Down-Regulation of *osRAFTIN1* Induces Male Sterility in the Transgenic Rice

All the transgenic lines were morphologically observed for the consequences of down-regulation of *osRAFTIN1* expression. The transgenic plants of the *osRAFTIN1* silenced- lines and control lines showed similar tillering and leafing ability, similar leaf size, similar internode elongation and similar overall plant sizes (**Figures 8A and 8B**). Panicle initiation in all the lines occurred approximately 90-100 days post transplanting. All the lines had the similar unifloral spikelet subtended by two paleae lying between two glumes. All the flowers shared the same color, size and shape, and had apparently similar

gynoecium with an ovary and two feathery stigmas. The androecium in the flower of all lines had six stamens and each of them had a filament and an anther. The mature panicle of the *osRAFTINI*-silenced lines, however, was about 15-20% longer than that of the control lines (**Figures 8C and 8D**). The spikelet of the *osRAFTINI*-silenced lines did not open and no anthesis took place at maturity even over long extended time (6 weeks) after the emergence of the panicle. No seeds or only a few seeds per plant (less than 1 seed per panicle) were produced in the *osRAFTINI*-silenced lines, in contrast to more than 10 seeds per panicle in the control lines (**Figures 8C, 8D, 8E, 8F, 8G and 8H**). The palea in the *osRAFTINI*-silenced lines kept green and did not turn yellow over an extended time (**Figures 8E and 8F**). The *osRAFTINI*-silenced lines were crossed with pollen from a control line to test if the female organ was impaired. The successful production of hybrid seeds confirmed their normal ovary viability and function (**Figure 8I**). Therefore, the sterility in the *osRAFTINI*-silenced lines arose from the male organ.

Accordingly, the anther in the *osRAFTINI*-silenced lines was subjected to scanning electron microscopy. In the *osRAFTINI*-silenced line, the mature anther was mostly malformed, non-dehiscent and 10-15% smaller in length (**Figures 9A and 9D**) and almost all the pollen grains were abnormal, showing signs of abortion (**Figures 9B and 9E**). But there was no detectable difference in the pollen surfaces (**Figures 9C and 9F**). In a germination assay, the germination rate of pollens in the *osRAFTINI*-silenced lines was 4.7% while the germination rate of pollens in the control lines was 74.3%. Thus, in the *osRAFTINI* down-regulated lines (male sterile lines), vegetative growth and flower development were apparently normal prior to anthesis and the poor fertility was due to the abnormal development of the anther resulting from down-regulation of *osRAFTINI* expression.

Transmission electron microscopy was applied to further examine anther development. In all the lines, the Ubisch body, the orbicular wall, and the exinewall were evident with no distinct difference (**Figures 10A and 10B**). In the control lines, the mature anther contained many round pollen grains rich in starch granules and other contents inside (**Figure 10A**). In contrast, the anther of the *osRAFTINI*-silenced lines contained mostly aborted, flat pollen grains with very little contents inside (**Figure 10B**). The tapetum degeneration underwent well in the control lines with little tapetal remains, whereas in the *osRAFTINI*-silenced lines, the tapetal degeneration was arrested apparently at the vacuolated microspore stage, leaving the half degenerated tapetum about 4 μ m in

thickness as clearly evident at the very mature stage when the endothecium wall was thickened (Figures 10C and 10D).

The cereal grains constituting more than 60% of total global agricultural production offer the major portion of the human diet. Understanding the molecular biology of anther development in cereal crops is of great importance for crop improvement. Based on studies in some model plants such as *Arabidopsis* and tobacco, it is known that the anther development is governed by a large number of genes with spatio-temporal expression specificity (Goldberg et al., 1993, Koltunow et al., 1990). It is estimated that in *Arabidopsis* there are approximately 3500 genes (13.7% of total predicted genes) which are specifically expressed in the anther and not in other floral and vegetative tissues (Sanders et al., 1999). Cereals have relatively larger genome sizes (hexaploid wheat 16000 Mb; rice 420-466 Mb) with more genes (rice: 32,000 to 55615 genes) (Goff et al., 2002, Yu et al., 2002) in comparison with *Arabidopsis* (125 Mb, 25498 genes) (The *Arabidopsis* Genome Initiative, 2000). Thus, it can be deduced that cereal anthers recruit more genes to execute microsporogenesis. Moreover, considering more than 50% rice genes do not have homologues in *Arabidopsis*, a considerable portion of anther-specific genes are probably unique to cereals, that may account for the morphologic and metabolic difference in the anther development between cereals and *Arabidopsis*. In this study we isolated the *RAFTIN1* group of anther-tapetum specific genes, *taRAFTIN1a*, *taRAFTIN1b* and *taRAFTIN1d* in wheat and *osRAFTIN1* in rice, that are apparently unique to grasses and not present in *Arabidopsis* or other eudicots. The localization of the RAFTIN protein onto the Ubisch body and the microspore exinewall raises the possibility that this protein is involved in transport of some macromolecules or their derivatives produced in the tapetum of cereals but not of *Arabidopsis*. Thus, study on *RAFTIN1* assists in understanding the molecular genetics underlying the metabolic difference in anther development between cereals and *Arabidopsis*.

The Ubisch Body-Mediated Transport of RAFTIN1 and the Function of RAFTIN1

Ubisch bodies, or orbicules are the minute sporopollenin particles comprising acidic and neutral polysaccharides, proteins and unsaturated lipids that line the inner surface of most secretory tapeta (El-Ghazaly & Jensen, 1986, El-Ghazaly & Jensen, 1987, Huysmans et al., 1998, Suarez-Cervera et al., 1995). Although discovered more than a century ago, Ubisch bodies, in terms of their structure, origin, development and function, remain mysterious. Based on cytological studies mainly through electron microscopy, it is

believed that Ubisch bodies originate from so-called "grey bodies" or "globular bodies" (pro-orbicule; pro-Ubisch bodies) derived from endoplasmic reticulum in the cytoplasm of the tapetum as early as the meiosis and tetrad stages (El-Ghazaly & Jensen, 1986, El-Ghazaly & Jensen, 1987, Huysmans et al., 1998, Suarez-Cervera et al., 1995).

5 Approaching the plasma membrane, pro-Ubisch bodies are bound by membranes (El-Ghazaly & Jensen, 1986). At the free microspore stage, pro-Ubisch bodies, upon fusion of the plasma membranes, are released from but connected with the plasma membrane by a layer of microfibril. Along with the coating and accumulation of sporopollenin on the surface of pro-Ubisch bodies, they mature to Ubisch bodies. At the same time,
10 sporopollenin is deposited between and beneath Ubisch bodies to form an orbicular wall (El-Ghazaly & Jensen, 1986). In the previous studies, functions hypothesized for the Ubisch body include transport of sporopollenin, temporary packing of sensitive material for transport through locular sap, by-products of tapetal cell metabolism, association with pollen dispersal, degradation of tapetal cells, and prevention against osmosis and collapse
15 of developing microspores (Huysmans et al., 1998).

In this study, we found that RAFTIN1 proteins were localized to the tapetum, the Ubisch body and the microspore exinewall, but not on the orbicular walls and inside microspore. Moreover, *RAFTIN1* transcript was only evident in the tapetum. Since Ubisch bodies are physically located between the plasma membrane of the tapetum and the
20 exinewall of microspore, it is therefore conceivable that through the Ubisch body, RAFTIN1 is transported from the tapetum to microspore. RAFTIN1 proteins were not detected until the microspore stage, suggesting that the deposition of RAFTIN1 into the Ubisch body and further onto the microspore exinewall is probably concurrent with that of sporopollenin. Silencing *RAFTIN1* expression did not disrupt or discernibly change the
25 structure of Ubisch bodies and the microspore exinewall, indicating RAFTIN1 is not required for building-up the basic skeleton of the Ubisch body and the microspore exinewall. However, we found that in the *RAFTIN1*-silenced lines, the tapetal degeneration was clearly retarded, and microspore was smaller and contained much less contents. One possible explanation is that RAFTIN1 is required for the degradation of
30 tapetal cells and the failure to such degradation terminates the nutrient and metabolite release from the tapetum to the locule leading developing microspore starving to abortion. However, this assumption is not in agreement with the following findings: RAFTIN1 proteins, though synthesized in the tapetum, are transported outside of the tapetum;

RAFTIN1 does not have any known domains for such a possible function; and over-expression of *RAFTIN1* does not induce cell degradation in a model plant. Therefore, *RAFTIN1* is not likely a candidate protein that involves the programmed cell death of the tapetum. Alternatively, *RAFTIN1* probably directly or indirectly regulates transport of certain metabolites which are rich in the tapetum of cereal crops, such as acid polysaccharides, neutral polysaccharides or their derivatives. Cereal pollen grains accumulate large amounts of starch granules (this study; (Bedinger, 1992) (Zhang et al., 2001). During microspore development, carbohydrate metabolism takes place very actively in the tapetum and in the surrounding supportive tissues as well as in microspores. In wheat, substantial amounts of acidic polysaccharides, proteins, neutral polysaccharides and to a lesser extent unsaturated lipids are found in the Ubisch body and the microspore exinewall over a long period during microspore development (El-Ghazaly & Jensen, 1987). Disturbance in carbohydrate metabolism induces male sterility (Dorion et al., 1996, Lalonde et al., 1997, Zhang et al., 2001). Interestingly, in crucifers and other entomophilous species where no *RAFTIN1*-like ESTs or genes have been documented, it is lipids not carbohydrates that are major products of the tapetum (Piffanelli et al., 1997), implying a difference between major tapetal metabolism of cereals and that of entomophilous species. The *RAFTIN1*-less Ubisch body and microspore exinewall probably is incompetent to transport these metabolites from the degenerating tapetum to microspores, which disconnects the nutrient supply-pipe to microspores leading to pollen abortion and, in turn, slows down the tapetal degeneration.

Molecular Hallmarks of BURP Domain-Containing Proteins and Their Multi-Functional Roles

The C-terminal moiety of *RAFTIN1* shares extensive sequence homology with a BURP domain that has been found only in the plant kingdom. These include *RD22* (a gene responsive to dehydration stress, high salt or ABA induction) from *Arabidopsis* (Yamaguchi-Shinozaki & Shinozaki, 1993), *A2-134* (or *ASG-1*) in the developing embryo of apomictic guinea grass (*Panicum maximum*) (Chen et al., 1999), the β subunit of polygalacturonase isoenzyme 1 (*PGI β*) in the developing fruit of tomato (Zheng et al., 1992), *USP* showing seed-specific in fava bean (*Vicia faba*) (Bäumlein et al., 1991), *BNM2* (a gene expressed during the induction of microspore embryogenesis) in rape (*Brassica napus*) (Hattori et al., 1998), *ADR6* (an auxin down-regulated gene) in soybean (Datta et al., 1993), *SALI 3-2* (an aluminum up-regulated gene) from soybean roots

(Ragland & Soliman, 1997) and some ESTs from *Prunus persica* (GenBank accession no: AAL26909), from cotton fiber cells (GenBank accession no: AAL67991), and from soybean seed coats (GenBank accession no: AAL76058; gene named *SCB1*). Thus, in spite of highly conserved primary structural feature within their C-terminal BURP

5 domains, the BURP domain proteins are expressed in divergent tissues and under various conditions. Nevertheless, these proteins are expressed in the tissues either under stress (chemicals, ABA, high salts, dehydration, auxin-down regulation), or during development (the anther, fruit and seed), which are undergoing active biodegradation and biosynthesis metabolism and probably intercellular metabolite movement. Alignments of the BURP
10 domains confirm the previous finding that there is a consensus sequence, CHX₁₀CHX₂₅₋₂₇CHX₂₄₋₂₅CH containing four repeated cysteine-histidine (CH) motifs located at C-termini (Hattori et al., 1998). These conserved motifs are the hallmarks of the BURP domains and are probably involved in the formation of disulfide bond intramolecularly for the proper protein folding or more likely intermolecularly for anchoring of the protein onto specific
15 sites of the cell wall. This later notion is supported by observations that all the BURP domain proteins are only found in the plant kingdom and that the two only characterized ones, *i.e.*, *RAFTIN1* (this study) and PG1 β (Zheng et al., 1992) have been subcellularly localized into the Ubisch body and the microspore exinewall, and the cell wall, respectively. The N-terminal moiety of the BURP domain proteins is highly divergent.

20 This divergence may reflect their multi-functionality. Thus, PG1 β , a 69 kDa non-catalytic fruit-specific cell wall glycoprotein that is proposed to play a role in the localization, immobilization or activation of the polygalacturonase enzyme complex within the cell wall, may use its C-terminal BURP domain for cell wall attachment, and its N-terminal domains for association with the catalytic subunit and regulation of its activity. In contrast,
25 *RAFTIN1* is anchored to the Ubisch body and the microspore exinewall by its BURP domain and its N-terminal moiety regulates metabolite transport. Further characterization of other BURP domain proteins will assist in unraveling the functional mystery of the BURP domain proteins, perhaps the key role of the biological process in anther development.

30 The presence of *RAFTIN* genes has been demonstrated in a number of plant species, especially those belonging to monocots. Thus, it appears that this gene is likely to be present in other monocots including other cereals and grasses. Silencing or knocking-out of *RAFTIN* will find utility in breeding programs where male sterile lines are required.

Furthermore, this technology can be used to prevent flow of transgenic pollens from elite transgenic lines, e.g., herbicide resistance lines, to wild plant species by silencing or knocking-out *RAFTIN* genes.

5 METHODS

Plant Materials

Hexaploid spring wheat (*Triticum aestivum* L. cv. *AC Karma*, AABBDD), tetraploid wheat (*T. turgidum* L. cv. *Sceptre*, AABB), two diploid wheat species (*T. urartu*, ssp. *nigrum*, AA; *T. tauschii* L. china, DD), were obtained as described previously (Wang et al., 2002) and rice (*Oryza sativa* L. japonica var. *nipponbare*) was obtained from National Institute of Agrobiological Resources, Japan, and were grown in a greenhouse.

Isolation of *taRAFTIN1a*, *taRAFTIN1b* and *osRAFTIN1* cDNAs, and Their Corresponding Genomic Sequences

A cDNA clone (A71) was obtained from a wheat anther cDNA library constructed previously (Wang et al., 2002). As BLASTN or BLASTX searches of 5' sequence of A71 against the *Arabidopsis* genome did not identify any A71-like sequence, the insert of clone A71 was completely sequenced. The 5' region of the predicted open reading frame (ORF) was amplified by PCR with primers OL3044 (5'TGCCACACTCGCCATTG3') (SEQ ID NO: 1) and OL3045 (5'TTTCCAGCGAGGCTGCT3') (SEQ ID NO: 2). The resulting 346 bp DNA fragment was used as a probe to screen ~ 500,000 clones of the anther cDNA library. The phagemids from the 26 positive plaques, excised *in vivo* using the ExAssist/SOLR system (Stratagene), were sequenced. Based on sequences, these cDNAs were placed into two groups, and the longest of each group was named *taRAFTIN1a* and *taRAFTIN1b* accordingly.

The entire *osRAFTIN1* ORF was cloned by RT-PCR using primers OL4382 (5'CGGGGTACCGAACGCTTCCATGGCGCGCT3' (SEQ ID NO: 3), *KpnI* site underlined, start codon ATG in italic) and OL4383 (5'GCTCTAGAGCTTCTACGCCCGTCGAGCTC3' (SEQ ID NO: 4), *XbaI* site underlined, the codon in italic is complementary to the stop codon TAG). The cDNA was directionally cloned into the *KpnI*-*XbaI* sites of plasmid pBluescript SK (Stratagene).

The genomic counterparts of *taRAFTIN1a*, *taRAFTIN1b* and *osRAFTIN1* were obtained by PCR of the genomic DNA, cloned into the T/A vector (Invitrogen, Carlsbad, CA, USA) and sequenced.

Oligonucleotide Synthesis, DNA Sequencing and Sequence Analysis

Oligonucleotide synthesis and DNA sequencing were carried out by the DNA Technology Unit of the Plant Biotechnology Institute. DNA sequences were assembled and analyzed using Lasergene software (DNASTAR Inc., Madison, WI, USA), FASTA (www.ebi.ac.uk/fasta3/), BLAST 2 and BLAST (www.ncbi.nlm.nih.gov).

DNA and RNA Gel Blot Analysis

Genomic DNA was isolated from leaves as described (Wang et al., 2002). DNA was digested with appropriate restriction enzymes, fractionated by electrophoresis on a 0.8% agarose gel, and transferred onto a Hybond N⁺ membrane (Amersham, Baie d'Urfe, Quebec, Canada). The methods for nucleic acid isolation, blotting, ³²P-labeling of probes and hybridization were as described (Wang et al., 2002). The entire ORFs of *taRAFTIN1a* and *osRAFTIN1* retrieved by PCR were used as probes for hybridization. Hybridization and subsequent washing conditions were essentially as described (Wang et al., 2002).

RT-PCR Analysis of *RAFTIN1* Gene Expression

First strand cDNA was generated in a 20 µl reaction containing 5 µg of total RNA isolated from appropriate wheat/rice tissues, 0.5 µg oligo (dT)₁₈, 20 units of SUPERScriptTM II RNase H⁻ Reverse Transcriptase (Invitrogen) according to the supplier's instruction. One hundred-fifty ng RNA-derived cDNA was used for a 100-µl PCR reaction in the presence of 10 units of Taq DNA polymerase (Amersham). Primers OL3044 and OL3073 (5'CTCCATGTCCACCATGTA3') (SEQ ID NO: 5) were used for amplification of an 875-bp fragment at the 5' coding region of wheat *taRAFTIN1a* cDNA, whereas primers OL3148 (5'CGACGTATTTGTCGTAGT3') (SEQ ID NO: 6) and OL3815 (5'TCTCGAACGCTTCCATG3') (SEQ ID NO: 7) were targeted at a 441-bp cDNA immediately from the putative start codon of rice *osRAFTIN1*. Primers OL4556 (5'TCGAGCTCGTCGCCGTCA3') (SEQ ID NO: 8) and OL4557 (5'GCAGCACCAGTGCTGCTG3') (SEQ ID NO: 9) binding to cDNA of a house-keeping gene, *glyceraldehyde-3-phosphate dehydrogenase* (GAPDH) were used as a control. All PCR was carried out with a Techne Genius thermocycler (Duxford, Cambridge, UK) as indicated: 35 cycles of 94°C, 30 sec; 56°C, 30 sec; and 72°C, 1 min; finally a 10-min extension at 72°C. Five µl of the reaction was used for agarose gel analysis.

Production and Purification of RAFTIN1 Polyclonal Antibodies

The 5' region of *taRAFTIN1a* ORF was amplified by PCR using a *Bam*HI site-containing 5' primer (5'CGGGATCCGCCTTCCTCGTCGC3' (SEQ ID NO: 10), *Bam*HI site underlined) and an *Eco*RI site-containing 3' primer

- 5 (5'GGAATTCTCACGCCGGCGAGCGATT3' (SEQ ID NO: 11), *Eco*RI site underlined) and cloned in-frame into the *Bam*HI-*Eco*RI sites of plasmid pTrxFus (Invitrogen, Carlsbad, CA, USA) to yield plasmid pAMWthio-A71. The fusion protein produced in *E. coli* strain GI724 (Invitrogen) hosting plasmid pAMWthio-A71 was purified (Wang & Sanfacon, 2000) and used for immunizing rabbits (by the staff at Veterinary Infectious
10 Diseases Organization, University of Saskatchewan, Saskatoon, Canada). Antiserum IgG was initially purified as described (Wang et al., 1999) and further purified using Affi-Gel 10 Gel (Bio-RAD, Mississauga, Ontario, Canada) following supplier's instructions.

Protein Extraction, Subcellular Localization and Western Blotting Analysis

- For total protein extraction, different wheat tissues (200 mg each) were
15 homogenized with 1 ml extraction buffer (50 mM Tris-HCl, pH 8.0, 150 mM NaCl, 1% Triton X-100, 1% sodium deoxycholate, 10 mM β -mercaptoethanol, 1 mM phenylmethylsulfonyl fluoride (PMSF) and 1 mM ethylenediaminetetraacetic acid (EDTA)). The homogenate was centrifuged at 10,000 xg at 4 °C for 10 min. The supernatant was used for protein blotting analysis.

- 20 For protein subcellular location, 1 g developing wheat anthers were homogenized with 2 ml PSL buffer (protein subcellular localization buffer: 100 mM Tris-HCl, pH 7.5, 5 mM MgCl₂, 10 mM KCl, 10% glycerol, 0.4 M sucrose, 10 mM β -mercaptoethanol, and 1 mM PMSF) on ice. The homogenate was filtered through an 80 mesh nylon cloth. The insoluble materials containing cell wall debris, after washing 3 times with PSL buffer plus
25 1% Triton X-100, were obtained as the cell wall fraction. The filtrate was centrifuged at 1,000xg for 10 min at 4 °C. The pellet rich in nucleus was called the nucleus fraction. The supernatant was centrifuged at 30,000xg for 30 min at 4 °C. The resulting pellet mainly containing membranous materials was the membrane-binding fraction and the supernatant containing the soluble protein was the cytosol fraction. All fractions were diluted to 2 ml
30 with PSL buffer. Equal volume of each fraction was used for western blot analysis.

For immunoblot analysis, proteins were separated by denaturing sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) (Laemmli, 1970) and transferred onto a polyvinylidene fluoride (PVDF) membrane using a Bio-Rad miniblotter at 125 V

for 2 hr in a Tris/MeOH/glycine buffer (25 mM Tris-HCl, pH 8.3, 20% (v/v) methanol, 192 mM glycine). The membrane was incubated in a blocking solution containing 3% bovine serum albumin (BSA) in TBS buffer (20 mM Tris-HCl, pH 7.5, 500 mM NaCl) with gentle agitation for 30 min, and then washed twice with TBS buffer for 5 min each. After incubation with the first antibodies in incubation buffer (1% BSA and 0.05% Tween-20 in TBS) for 1 hr, the filter was washed with washing buffer (0.05% Tween-20 in TBS) for 5 min three times. Secondary antibodies conjugated with alkaline phosphatase (goat anti-rabbit IgG; Bio-Rad) (1:2000 diluted with incubation buffer) were incubated with the filter for 1 hr. The filter was washed three times with washing buffer and twice with alkaline phosphatase buffer (AP buffer: 100 mM Tris-HCl, pH 9.5, 100 mM NaCl, 50 mM MgCl₂). The antibody-protein complex was visualized with enzyme substrates, BCIP (bromochloroindolyl phosphate; Bio-Rad) and NBT (nitro blue tetrazolium; Bio-Rad) in AP buffer. The reaction was stopped by addition of excess stop buffer (20 mM Tris-HCl, pH 7.5, 5 mM EDTA). All steps were performed at RT except otherwise stated.

***In situ* RNA Hybridization and Immunocytochemistry**

Rice and hexaploid wheat flower sections, *in situ* RNA hybridization and immunocytochemistry were performed as described (Wan et al., 2002, Wang et al., 2002).

Promoter isolation

The upstream regulatory regions of *taRAFTIN1a* and *taRAFTIN1b* coding regions were isolated from the hexaploid wheat cultivar Karma (genetic complements: AABBDD) using a Universal GenomeWalkerTM Kit (Clontech, Palo Alto, CA). Two nested reverse primers, OL3070 (5'TCCAGCCTGAACCGCGACCAGGGTGGT3') (SEQ ID NO: 12) and OL3071 (5'GTGGTGGCGAGGAGGGCGACGAGGAA3') (SEQ ID NO: 13) were used for the first and second PCR. The resulting two fragments of 1.7 kb for *taRAFTIN1a* and 2.1 kb for *taRAFTIN1b* were cloned into a T/A vector (Invitrogen). The inserts were completely sequenced.

A DNA fragment 1458 bp upstream of the predicted start codon for *osRAFTIN1* ORF was retrieved directly by PCR of the genomic DNA using primers OL3079 (5'CGAAGGACTCTGGT3') (SEQ ID NO: 14) and OL3080 (5'CATGGAAGCGTTTCGAGA3') (SEQ ID NO: 15) and cloned into a T/A vector (Invitrogen).

Transformation Vector Construction and Plant Transformation

For promoter analysis in tobacco and Arabidopsis, cloned promoters were retrieved using appropriate sets of primers: OL3861

(5'CCCAAGCTTCTGTCGATGGCGCTCTGT3' (SEQ ID NO: 16), *Hind*III site

underlined) and OL3862 (5'CGGGATCCGATGTGCGCTAGGTGAGA3' (SEQ ID NO: 17), *Bam*HI site underlined) for amplification of *taRAFTIN1a* upstream regulatory region (1716 nt upstream of start codon), OL3863

(5'CAAGCTTCTAGACTTGTTGAGTGCCACACT3' (SEQ ID NO: 18), *Hind*III site underlined) and OL3862 for *taRAFTIN1b* (2092 nt upstream of start codon), OL3142

(5'CCCAAGCTTTACCCACACGTCATGA (SEQ ID NO: 19), *Hind*III site underlined) and OL3143 (5'CGGGATCCCATGGAAGCGTTCGAGA3' (SEQ ID NO: 20), *Bam*HI site underlined) for *osRAFTIN1* (1309 nt upstream of start codon). The PCR products were restricted and ligated into the *Hind*III-*Bam*HI sites of the modified plasmid pAMW477

(Wang et al., 2002) in which the *TAA1a* coding region had been replaced with a *GUS* gene from plasmid pRD420 (Datla et al., 1992) to generate plasmid pAMW484

(*taRAFTIN1a::GUS* chimeric gene), pAMW483 (*taRAFTIN1b::GUS*), and pAMW486

(*osRAFTIN1::GUS*). Genetic transformation of *Nicotiana tabacum* cv. Xanthi and

Arabidopsis thaliana ecotype Col-0 were essentially as described (Clough & Bent, 1998,

Wang et al., 2002). For promoter analysis in rice, the *Hind*III-*Kpn*I fragment of plasmids

pAMW484, pAMW483 and pAMW486 containing the *RAFTIN1::GUS* cassette were individually co-ligated with an *Hind*III-*Kpn*I fragment (35S::*hph* cassette) of plasmid pBShph (R. Datla, Plant Biotechnology Institute, National Research Council of Canada, Canada) to produce plasmids pAMW 499, pAMW494 and pAMW501, respectively.

ihpRNA intermediate clone pAMW487 containing *RAFTIN1a* promoter,

complementary sequence of *osRAFTIN1* cDNA (nt 159 from start codon to nt 415), its corresponding sense genomic DNA sequence with additional 114 nt of upstream genomic sequence including an 82 nt intron, and a 35S terminator (hereafter named *RAFTIN1::hp-osRAFTIN1*) was constructed by co-ligation of a *Bam*HI-*Spe*I restricted PCR fragment of

osRAFTIN1 cDNA amplified with OL3888 (5'CGGGATCCGACGTATTTGTCGTAGT3' (SEQ ID NO: 21), *Bam*HI site underlined) and OL3889

(5'GGACTAGTCAGCTTCGTCGTCGGCA3' (SEQ ID NO: 22), *Spe*I site underlined)

and an *Xba*I-*Eco*RI restricted PCR fragment of *osRAFTIN1* genomic DNA with OL3886 (5'GCTCTAGACGCCTTCCTCCGCCT3' (SEQ ID NO: 23), *Xba*I site underlined) and

OL3887 (5'GGAATTTCGACGTATTTGTCGTAGT3' (SEQ ID NO: 24), *Eco*RI site underlined) into *Bam*HI-*Eco*RI sites of plasmid pAMW484. Similar strategy was used to make clone pAMW488 containing *taRAFTIN1b::hp-osRAFTIN1*, clone pAMW489 containing *osRAFTIN1::hp-osRAFTIN1*, and clone pAMW503 (*35S::hp-osRAFTIN1*).

5 Clone pAMW491 containing *taRAFTIN1b* promoter, complementary sequence of *taRAFTIN1a* cDNA (nt 181 from start codon to nt 544), its corresponding sense genomic sequence with additional 148 nt of upstream genomic sequence including a 99 nt intron), and a 35S terminator (hereafter named *taRAFTIN1b::hp-taRAFTIN1a*) was created by coligation of a *Bam*HI-*Spe*I restricted PCR fragment of *taRAFTIN1a* cDNA amplified
10 with OL3892 (5'CGGGATCCTGGGAGCCTCTTGCCGA3' (SEQ ID NO: 25), *Bam*HI site underlined) and OL3893 (5'GGACTAGTCACAGAAGCCACCAGCT3' (SEQ ID NO: 26), *Spe*I site underlined) and an *Xba*I-*Eco*RI restricted PCR fragment of *taRAFTIN1a* genomic DNA with OL3890 (5'GCTCTAGACGCCGTTCTCCGCCT3' (SEQ ID NO: 27), *Xba*I site underlined) and OL3891

15 (5'GGAATTCTGGGAGCCTGTTGCCGA3' (SEQ ID NO: 28), *Eco*RI site underlined) into *Bam*HI-*Eco*RI sites of plasmid pAMW485. Similar strategy was employed to generate subclones pAMW492 containing *osRAFTIN1::hp-taRAFTIN1a* and pAMW504 containing *35S::hp-taRAFTIN1a*.

Subclone pAMW502 containing *taRAFTIN1a::hp-taRAFTIN1a* was constructed
20 by insertion of the small fragment from *Bam*HI-*Kpn*I double digested pAMW491 into the corresponding sites of pAMW484. The small fragment of clones pAMW487 (3.1 kb), pAMW488 (2.9 kb), pAMW 489, pAMW491 (2.8 kb), pAMW492 (3.0 kb), pAMW502 (3.0 kb), pAMW503 (2.8 kb) and pAMW504 (2.9 kb) double-digested with *Hind*III-*Kpn*I was ligated with the small 1.7 kb *Hind*III-*Eco*RI fragment (*35S::hph* cassette) of plasmid
25 pBShph and plasmid pHS723 (Huang et al., 2000) digested with *Kpn*I and *Eco*RI to obtain ihpRNA transformation vectors pAMW495, pAMW496, pAMW497, pAMW498, pAMW500, pAMW506, pAMW507 and pAMW508, respectively. Rice transformation was carried out following the published protocol (Chen et al., 1998).

Pollen Viability Assay

30 Pollen from mature anthers of the *osRAFTIN1*-silenced lines or the control lines was stained with 1% aniline blue in lactophenol. The viability of pollen was calculated based on over 300 pollen grains.

Scanning Electron Microscopy (SEM), Transmission Electron Microscopy (TEM) and Immunoelectron Microscopy (IM)

For SEM, mature anthers and rice pollen grains were mounted on aluminum stubs by a piece of double-sided tape (Canemco, St. Laurent, Quebec, Canada) and were
5 sputtered with gold. The specimens were observed in a Phillip 505 scanning electron microscopy (Philips Electron Optics, Eindhoven, The Netherlands).

For TEM, rice spikelets were fixed in 3% glutaraldehyde in 0.025 M phosphate buffer, pH 6.8 overnight at 1 hr 4 °C and postfixed in 1% osmium tetroxide on ice for 8 hrs. After dehydration in a graded ethanol series, the samples were embedded in acrylic
10 resin (London Resin Company, Reading, Berkshire, UK). Ultra-thin sections (50-70 nm) were made using a Reichert Jung Ultracut E microtome (Leica, Vienna, Austria), and double-stained with 2% (w/v) uranyl acetate and 2.6% (w/v) lead citrate. The section was viewed and photographed with a Philips CM-10 transmission electron microscope (Philips Electron Optics).

For IM, developing anthers were fixed with 1.5% glutaraldehyde in 0.025 M phosphate buffer, pH 6.8 for 1 hr and then 3% glutaraldehyde in the same buffer for 3 hr at room temperature, rinsed with phosphate buffer at 4 °C overnight and dehydrated in a graded ethanol series. The fixed anthers were infiltrated with LR-white resin and polymerized with UV light. Sections (0.5 µm) were cut using a microtome (Reichert
20 Ultracut E, BEMF, Honolulu, Hawaii, USA) and mounted on a silicone rubber plate (Canemco Inc, St. Laurent, Quebec, Canada) on a 300-mesh carbocoated nickel grid. The section was incubated with blocking solution containing 1% BSA in PBS buffer (10.14 mM Na₂HPO₄, 1.76 mM KH₂PO₄, pH 7.4, 136.9 mM NaCl, 2.69 mM KCl) for 30 min, followed with one hr incubation with the column purified antibody in the blocking
25 solution (0.01 mg/ml). After washing 3 times with PBS buffer for 15 min, the section was reacted with the gold-labeled goat anti-rabbit IgG (EMGAR15; British BioCell International, Cardiff, UK) (100-fold dilution with the blocking solution) for 1 hr. The section was washed with 3 changes of PBS buffer for 15 min and 4 changes of distilled water for 12 min. The grid were stained with 2% uranyl acetate for 20 min, washed 4
30 times with distilled water for 16 min and incubated with 0.3% lead citrate for 10 min, followed with 4 times of rinse with distilled water for 20 min. The section was viewed and photographed with a Philips 410 LS electron microscope (Philips Electron Optics). All the above steps were performed at RT except otherwise stated.

By way of examples, we have shown reduction of male fertility by silencing the expression of a *RAFTIN* gene in rice. It should be possible to enhance male fertility in plants by modulating appropriately sustained *RAFTIN* gene expression in a plant. For this purpose, a *RAFTIN* nucleotide sequence would be placed in sense orientation under the control of an anther-expressing promoter and using standard transformation vectors a plant would be transformed. The transformed cell is selected and grown into a plant and analyzed for male fertility at the time of flowering. The plant is found to have enhanced fertility.

10 REFERENCES

- Aarts, M. G. M., Hodge, R., Kalantidis, K., Florack, D., Wilson, Z. A., Mulligan, B., Stiekema, W. J., Scott, R. & Pereira, A. (1997). The Arabidopsis MALE STERILITY 2 protein shares similarity with reductases in elongation condensation complexes. *Plant Journal* 12, 615-623.
- 15 Bäumlein, H., Boerjan, W., Nagy, I., Bassünér, R., Van Montagu, M., D., I. & Wobus, U. (1991). A novel seed protein gene from *Vicia faba* is developmentally regulated in transgenic tobacco and Arabidopsis plants. *Molecular General Genetics* 225, 459-467.
- Bedinger, P. A. (1992). The remarkable biology of pollen. *Plant Cell* 4, 879-887.
- 20 Chen, L., Miyazaki, C., Kojima, A., Saito, A. & Adachi, T. (1999). Isolation and characterization of a gene expressed during early embryo sac development in apomictic guinea grass (*Panicum maximum*). *J. Plant Physiol.* 154, 55-62.
- Chen, L., Zhang, S., Beachy, R. N. & Fauquet, C. M. (1998). A protocol for consistent, large-scale production of fertile transgenic rice plants. *Plant Cell Report* 18: 25-31.
- 25 Plant Cell Report 18, 25-31.
- Clough, S. J. & Bent, A. F. (1998). Floral dip: a simplified method for *Agrobacterium*-mediated transformation of *Arabidopsis thaliana*. *The Plant Journal* 16, 735-743.
- D'Arcy, W. G. (1996). Anthers and stamens and what they do. In *The anther: form, function and phylogeny*, pp. 1-24. Edited by W. G. D'Arcy & R. C. Keating. Cambridge, UK: Cambridge University Press.
- 30 Datla, R. S. S., Hammerlindl, J. K., Panchuk, B., Pelcher, L. E. & Keller, W. (1992). Modified binary plant transformation vectors with the wild-type gene encoding NPTII. *Gene* 122, 383-384.

- Datta, N., LaFayette, P. R., Kroner, P. A., Nagao, R. T. & HKey, J. L. (1993). Isolation and characterization of three families of auxin down-regulated cDNA clones. *Plant Molecular Biology* 21, 859-869.
- De Block, M., Debrouwer, D. & Moens, T. (1997). The development of a nuclear male sterility system in wheat. Expression of the barnase gene under the control of tapetum specific promoters. *Theor Appl Genet* 95, 125-131.
- Dorion, S., Lalonde, S. & Saini, H. S. (1996). Induction of male sterility in wheat (*Triticum aestivum* L.) by meiotic-stage water deficit is preceded by a decline in invertase activity and changes in carbohydrate metabolism in anthers. *Plant Physiology* 111, 137-145.
- El-Ghazaly, G. & Jensen, W. A. (1986). Studies of the development of wheat (*Triticum aestivum*) pollen I. Formation of the pollen wall and Ubisch bodies. *Grana* 25, 1-29.
- El-Ghazaly, G. & Jensen, W. A. (1987). Development of wheat (*Triticum aestivum*) pollen. II Histochemical differentiation of wall and Ubisch bodies during development. *American Journal of Botany* 74, 1396-1418.
- Furness, C. A. & Rudall, P. J. (2001). The tapetum in basal angiosperms: early diversity. *International Journal of Plant Sciences* 162, 375-392.
- Goff, S. A. et al. (2002). A draft sequence of the rice genome (*Oryza sativa* L. ssp. japonica). *Science* 296, 92-100.
- Goldberg, R. B., Beals, T. P. & Sanders, P. M. (1993). Anther development: basic principles and practical applications. *Plant Cell* 5, 1217-1229.
- Hattori, J., Boutilier, K. A., van Lookeren Campagne, M. M. & Miki, B. L. (1998). A conserved BURP domain defines a novel group of plant proteins with unusual primary structures. *Molecular General Genetics* 259, 424-428.
- Huang, J., Hirji, R., Adam, L., Rozwadowski, K. L., Hammerlindl, J. K., Keller, W. A. & Selvaraj, G. (2000). Genetic engineering of glycinebetaine production toward enhancing stress tolerance in plants: metabolic limitations. *Plant Physiology* 122, 747-756.
- Huysmans, S., El-Ghazaly, G. & Smets, E. (1998). Orbicules in angiosperm: morphology, function, distribution, and relation with tapetum types. *Bot. Rev.* 64, 240-272.

- Koltunow, A. M., Truettner, J., Cox, K. H., Wallroth, M. & Goldberg, R. B. (1990). Different temporal and spatial gene expression patterns occur during anther development. *Plant Cell* 2, 1201-1224.
- Laemmli, U. K. (1970). Cleavage of structural proteins during assembly of the head of bacteriophage T4. *Nature (London)* 227, 680-685.
- Lalonde, S., Beebe, D. U. & Saini, S. (1997). Early signs of disruption of wheat anther development associated with the induction of male sterility by meiotic-stage water deficit. *Sexual Plant Reproduction* 10, 40-48.
- Mariani, C., Gossele, V., de Beuckeleer, M., de Block, M., Goldberg, R. B., de Greef, W. & Leemans, J. (1992). A chimaeric ribonuclease-inhibitor gene restores fertility to male sterile plants. *Nature* 357, 384-387.
- Mascarenhas, J. P. (1975). The biochemistry of angiosperm pollen development. *Bot Rev* 41, 259-314.
- Mascarenhas, J. P. (1990). Gene activity during pollen development. *Ann. Rev. Plant Physiol. Plant Mol. Biol.* 41, 317-338.
- McCormick, S. (1993). Male gametophyte development. *Plant Cell* 5, 1266-1275.
- Mizelle, M. B., Sethi, R., Ashton, M. E. & Jensen, W. A. (1989). Development of the pollen grain and tapetum of wheat (*Triticum aestivum*) in untreated plants and plants treated with chemical hybridizing agent RH00007. *Sex Plan Reprod* 2, 231-253.
- Piffanelli, P., Rogers, J. H. E. & Murphy, D. J. (1997). Intra and extracellular lipid composition and associated gene expression patterns during pollen development in *Brassica napus*. *Plant Journal* 11: 549-652.
- Ragland, M. & Soliman, K. M. (1997). Sali5-4a and Sali3-2, two genes induced by aluminum in soybean roots. *Plant Physiology* 114, 395.
- Sanders, P. M., Bui, A. Q., Weterings, K., McIntire, K. N., Hsu, Y.-C., Lee, P. Y., Truong, M. T., Beals, T. P. & Goldberg, R. B. (1999). Anther developmental defects in *Arabidopsis thaliana* male-sterile mutants. *Sex Plan Reprod* 11, 297-322.
- Shivanna, K. R., Cresti, M. & Ciampolini, F. (1997). Pollen development and pollen-pistil interaction. In *Pollen biotechnology for crop production and improvement*. Edited by K. R. Shivanna & V. K. Sawhney. Cambridge, United Kingdom: Cambridge University Press.

- Smith, N. A., Singh, S. P., Wang, M.-B., Stoutjesdijk, P. A., Green, A. G. & Waterhouse, P. M. (2000). Total silencing by intron-spliced hairpin RNAs. *Nature* 407, 319-320.
- Suarez-Cervera, M., Marquez, J. & Seoane-Camba, J. (1995). Pollen grain and Ubisch
5 body development in *Platanus acerifolia*. *Rev. Palaeobot. Palynol* 85, 63-84.
- The Arabidopsis Genome Initiative (2000). Analysis of the genome sequence of the flowering plant *Arabidopsis thaliana*. *Nature* 408, 796-815.
- Wan, L., Xia, Q., Qiu, X. & Selvaraj, G. (2002). Early stages of seed development in
10 *Brassica napus*: a seed coa-specific cysteine proteinase associated with programmed cell death of the inner integument. *Plant Journal* 30, 1-10.
- Wang, A., Carrier, K., Chisholm, J., Wieczorek, A., Huguenot, C. & Sanfacon, H. (1999). Proteolytic processing of tomato ringspot nepovirus 3C-like protease precursors: definition of the domains for the VPg, protease and putative RNA-dependent RNA polymerase. *Journal of General Virology* 80, 799-809.
- 15 Wang, A. & Sanfacon, H. (2000). Proteolytic processing at a novel cleavage site in the N-terminal region of the tomato ringspot nepovirus RNA-1-encoded polyprotein in vitro. *Journal of General Virology* 81, 2771-2781.
- Wang, A., Xia, Q., Xie, W., Dumonceaux, T., Zou, J., Datla, R. & Selvaraj, G. (2002). Male gametophyte development in bread wheat (*Triticum aestivum* L.): molecular,
20 cellular, and biochemical analyses of a sporophytic contribution to pollen wall ontogeny. *Plant Journal* 30, 613-623.
- Wiermann, R. & Gubatz, S. (1992). Pollen wall and sporopollenin. *Int. Rev. Cytol.* 140, 35-72.
- Yamaguchi-Shinozaki, K. & Shinozaki, K. (1993). The Plant hormone abscisic acid
25 mediates the drought-induced expression but not the seed-specific expression of rd22, a gene responsive to dehydration stress in *Arabidopsis thaliana*. *Molecular General Genetics* 238, 17-25.
- Yu, J. et al. (2002). A draft sequence of the rice genome (*Oryza sativa* L. ssp. indica). *Science* 296, 79-92.
- 30 Zhang, Y., Shewry, P. R., Jones, H., Barcelo, P., Lazzeri, P. A. & Halford, N. G. (2001). Expression of antisense SnRK1 protein kinase sequence causes abnormal pollen development and male sterility in transgenic barley. *Plant Journal* 28, 431-441.

Zheng, L., Heupel, R. C. & DellPenna, D. (1992). The β subunit of tomato fruit polygalacturonase isoenzyme 1: isolation, characterization, and identification of unique structural features. *Plant Cell* 4, 1147-1156.

CLAIMS

1. An isolated *RAFTIN* nucleotide sequence.
2. The isolated *RAFTIN* nucleotide sequence of claim 1 comprising a BURP domain
5 of two conserved phenylalanine residues (FF) at the N-terminus of the domain and a
consensus sequence of the formula $\text{CHX}_{10}\text{CHX}_{25-27}\text{CHX}_{24-25}\text{CH}$ wherein CH represents
each of four repeated cysteine-histidine (CH) motifs in the consensus sequence and X
represents any amino acid.
- 10 3. The isolated *RAFTIN* nucleotide sequence of claim 1 in the form of genomic DNA.
4. The isolated *RAFTIN* nucleotide sequence of claim 1 in the form of cDNA.
5. The isolated *RAFTIN* nucleotide sequence according to claim 1 comprising a
15 *RAFTIN* gene promoter sequence.
6. The isolated *RAFTIN* nucleotide sequence according to claim 1 comprising a
RAFTIN protein encoding sequence.
- 20 7. A transformation vector comprising a *RAFTIN* nucleotide sequence according to
claim 1.
8. A plant cell comprising the transformation vector of claim 7.
- 25 9. A plant comprising the transformation vector of claim 7.
10. A seed of a plant comprising the transformation vector of claim 7.
11. An RNA hairpin construct comprising a promoter operably linked to sense-
30 oriented *RAFTIN* nucleotide sequence, an intron and an antisense-oriented *RAFTIN*
nucleotide sequence.

12. A transformation vector comprising a sequence encoding the RNA hairpin construct of claim 11.

13. The transformation vector of claim 12 wherein said sequence encoding the RNA hairpin construct is operably linked to and under the control of a *RAFTIN* gene promoter sequence.

14. A method for producing a cell of a plant having male sterility or modulated male fertility which method comprises transforming said cell with the transformation vector of claim 12.

15. A method for producing a plant having male sterility or modulated male fertility which method comprises transforming a cell of said plant with the transformation vector of claim 12 and multiplying said cell to yield said plant.

16. A transformed plant cell comprising a sequence encoding the RNA hairpin construct of claim 11.

17. A male-sterile plant or seed thereof comprising transformed plant cells as defined in claim 16.

18. A method for producing a cell of a plant having enhanced male fertility which method comprises transforming said cell with the transformation vector of claim 7.

19. A method for producing a plant having enhanced male fertility which method comprises transforming said cell with the transformation vector of claim 7 and multiplying said cell to produce said plant.

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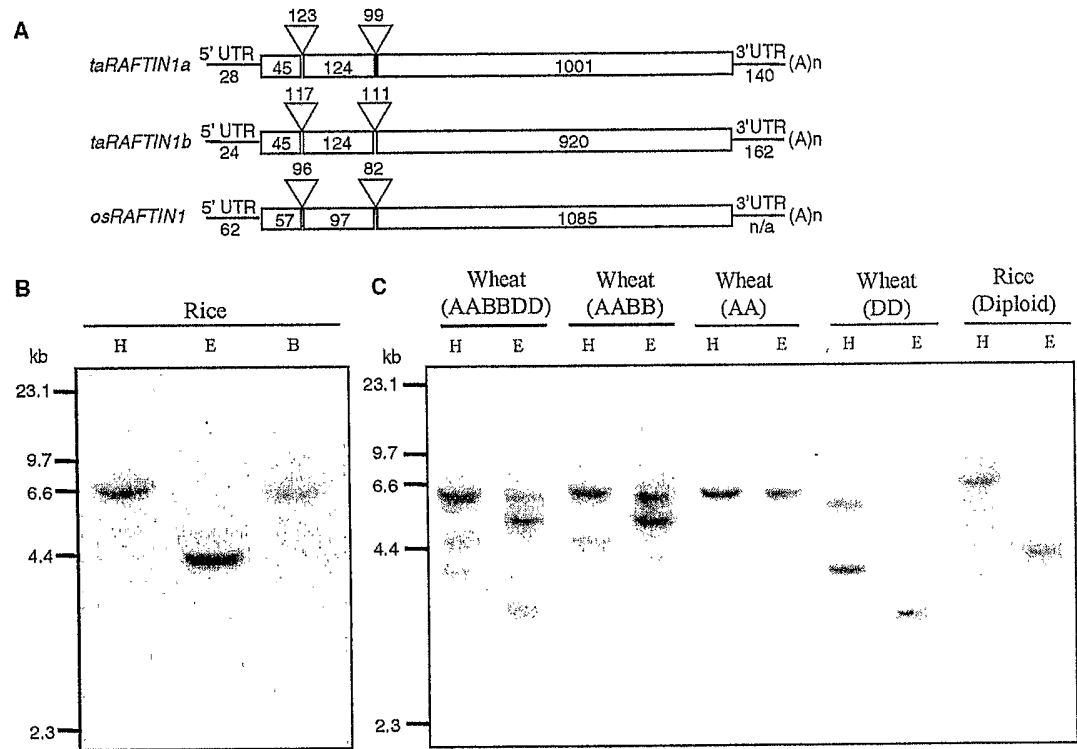


Figure 1

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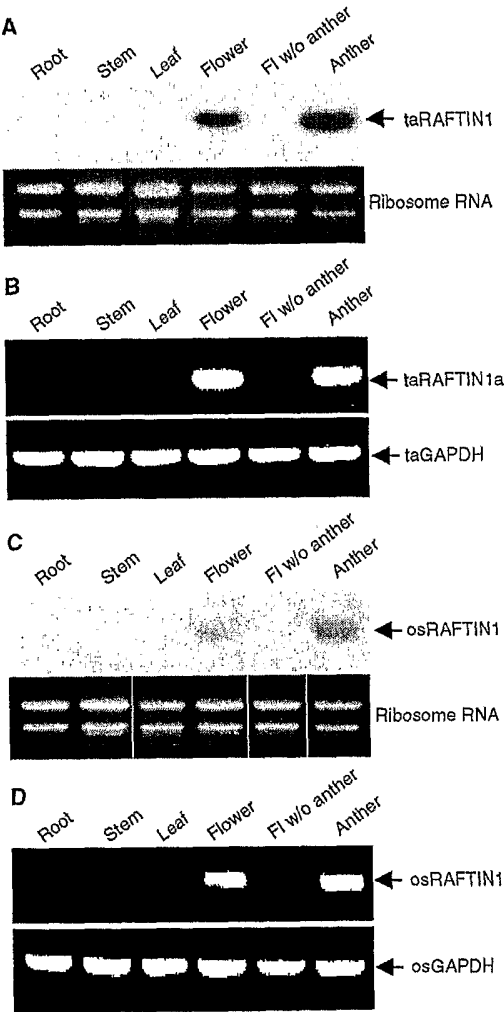


Figure 2 Aiming Wang *et al.* 2002

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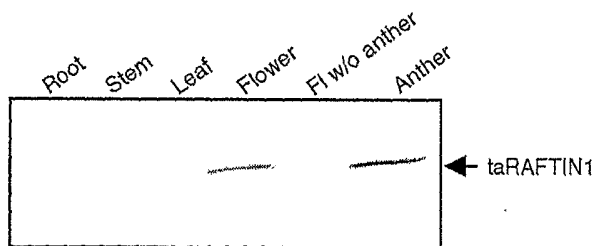


Figure 3 Aiming Wang *et al.* 2002

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A

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taRAFTIN1a 1  MARELVALLATTIVAVOAGGQLGHAAPATAEVFWRAVLPHSPLPDAVLRLLKQPAAGVELLTEATSFVRDAEDRAAPFD 78
taRAFTIN1b 1  -----S-----P-----AA----- 72
osRAFTIN1 1  -----AA--LVAVA-AA-VLS--D--S-----F-----RPDTSF-VGKA--AGAAAAGAARTGF-- 74

taRAFTIN1a 79  YRDYSRSPDPDEPSKSTGAAAASGARDFDYDDYSAAAAGGDKLRGAASGARAAAAADFDYDDYSGADKLRGATDAAAA 141
taRAFTIN1b 73  -----SS-----V-AA-----AAAAA-----N-----ER-----AAAA 114
osRAFTIN1 75  -T--RG-DSPTTA-GLDL-GDEGEPAF-G-A--AQGE--GGGA--A-EQVLAVDAG-N--K-V--R-L--GSSTAGGE 153

taRAFTIN1a 142  AAAAAAAEYKAPSSSLAGNGASMARGAGKAETTTVFHEEAVRVGKRLPFRFPFPAALGFLPRQVADSVPFTTAALP 212
taRAFTIN1b 115  AAAAAAA-----S-Y--A-----R--H----- 185
osRAFTIN1 154  NDDEPFGYD-----GSGTAATT--V-TGA-----E--Y--A-TS-----R-----I--A-- 233

taRAFTIN1a 213  GVLATFGVASDSATVASMEATLRACESPTIAGESKFCATSLALVERAMEVLGTRDIRPVTSTLPRAGAPLQTYTVRSVR 292
taRAFTIN1b 186  -I-----I-----T--P-----G-----VA-Q 265
osRAFTIN1 234  A--L-----P-T-EA-G-RE--T-W--L-----G-AA-----AALA--G--A-A--A-L 313

taRAFTIN1a 293  PVEGGPVFVACHDEAYPYTVYRCHTTGFSRAYMVDMEGAAARGGDAVTIATVCHTDTSLWNPEHVSFKLLGTPGGTTPV 369
taRAFTIN1b 266  -----T--AAA--A-----A----- 342
osRAFTIN1 314  ---AG--Q-----A--E--DGGGD--E--V--N--R-----S-- 392

taRAFTIN1a 370  CHLMPYGHIIWAKNVNRSPA 389
taRAFTIN1b 343  -----K----- 362
osRAFTIN1 393  -----V-----KS-T- 412

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B

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taRAFTIN1a 169  FFHEEAVRVGKRLPFRFPFPAALGFLPRQVADSVPFTTAALPGVLATFGVASDSATVASMEATLRACESPTIAGES 246
PG-bet 415  --R-KMLKS-TIMPAA-DIKDKMPKRS--VI-SKL--S-SKIAELKAI FHAGDE-QVEKMIGDA-SE--RAPS---T 492
RD22 176  --L-KDLVR--EMNV--NAEDGYGGKTA--GE-ET--GSEKFSET-KR-S-EAG-EEAEM-KK-IEE--ARKVSG-E 255
ASG-1 87  --N-HD-LE--TF-MY--SVTKA-K--Q-R-VOEI--S-R-ADI--L-HIPPG-SEAADVAT--GL-DAAAHGDVV 164
CFC1 124  --L-KDMHP-ATMSLH-TEN-BAAKSA--Y-T-QKI--SSDK--EIFNK-S-KPG-LK-EM-KN-IKE--Q-A-E--E 200
SCB1 92  --L--DL-A--IFNMK-VNN-AAA-TVPL--ISKQI--SEDKKKQ--ML--EAN-SNAKIAE--GL-QE-ATEG-R 169

taRAFTIN1a 247  KFCATSLALVERAMEVLGTRDIRPVTSTLPRAGAPLQTYTVARSVRPVEGGPVFVACHDEAYPYTVYRCHTTGFSRAY 324
PG-bet 493  -R-VN-A-DMIDEFATS--RNVAA-RT-EDTKGSNGNIMIGSVKGINGGKVTKSA-S--QTL--LL-Y--SVFKV-V- 569
RD22 256  -Y-----SM-DFSYSK--KIHV-A-STEVAKKN--M-K-KIAAAG-KKLSDDKA--V--KQK--FA-FY--KAMMTTV- 334
ASG-1 165  RA-V--PDDM-G-AA--SNMQVLAPS--TG-MS--P--AA-A-K--D-SDA--G--P-L--S-----SVQGTG- 241
CFC1 201  -Y-----SMIDYSISK--KV-AQA-STEVEKQATPM-K--IAAG-QKMTDDKA--V--KQN-A-A-FY--KSETT-- 276
SCB1 170  -H-----SM-DFVSA--KANVGAFSTEKERETESAGKFV-VKNG--KLGGDDHVAIA--PMS--V-FG--LVPR-SG- 246

taRAFTIN1a 325  MVDMEGARGAGDAVTIATVCHTDTSLWNPEHVSFKLLGTPGGTTPVCHLMPYGHIIW 380
PG-bet 570  EA-ILDPNSKVKINHGVAI--V--S-G-S-GA-VA--SG--KIE--WIFENDMTW 626
RD22 335  A-PL--AAEN-MRAKAVA--KN--A--N-LA--V-KV--TV--FL-ET-VV- 389
ASG-1 242  VME-QSSY-N-G-LKLVA--RN-TS-D-----V-AS--L-I--FV--V-F 298
CFC1 277  --PL--AAD-TKAKAVA--A--K-LA-QV-KVE--TI--FL-RD--V- 331
SCB1 247  L-RLK-AAED-VR-KAVVA--R--K-DHN-GA--V-NL--NGT--VFTE-NLL- 301

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Top

Figure 4 Aiming Wang *et al.* 2002

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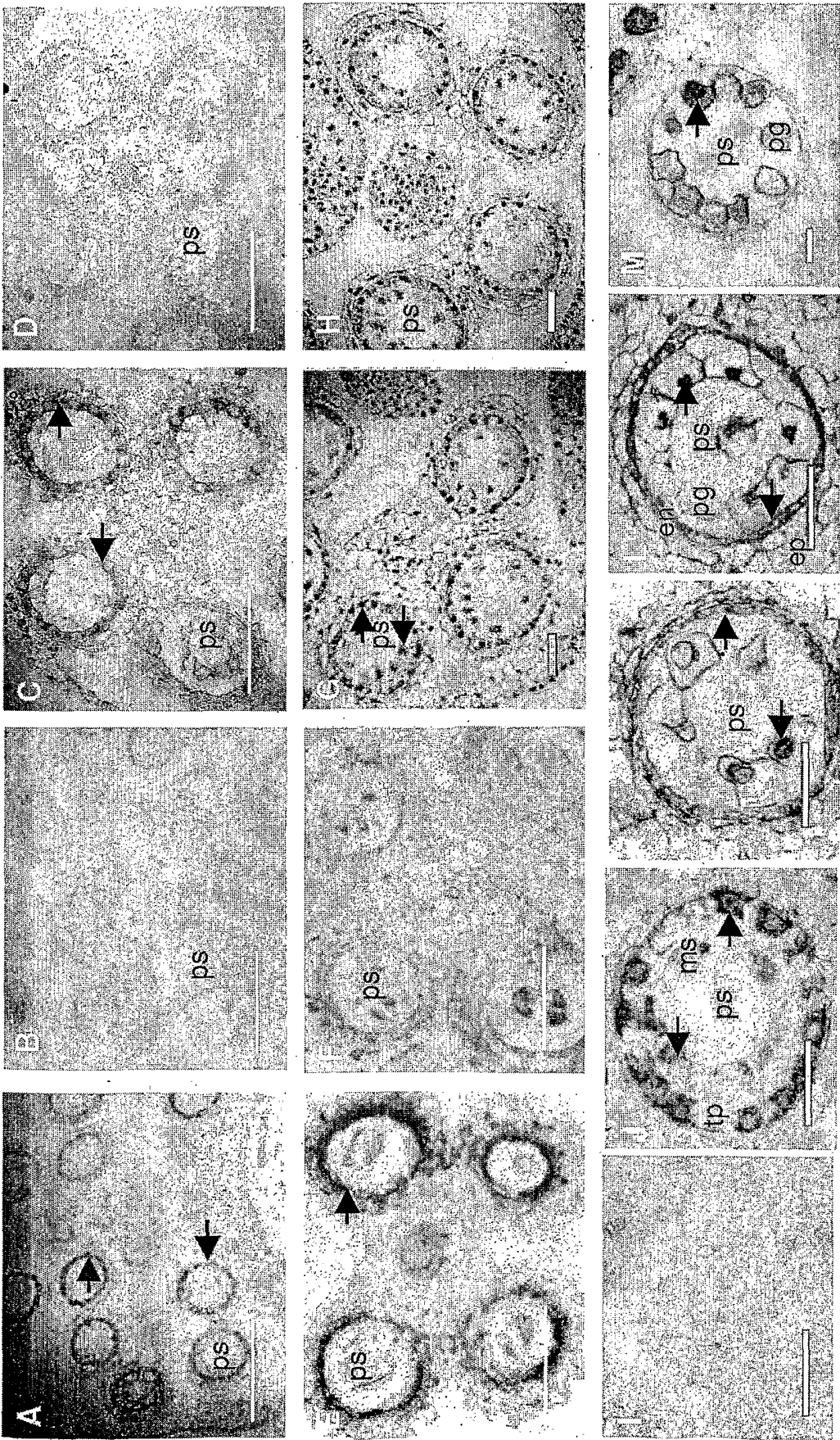


Figure 5

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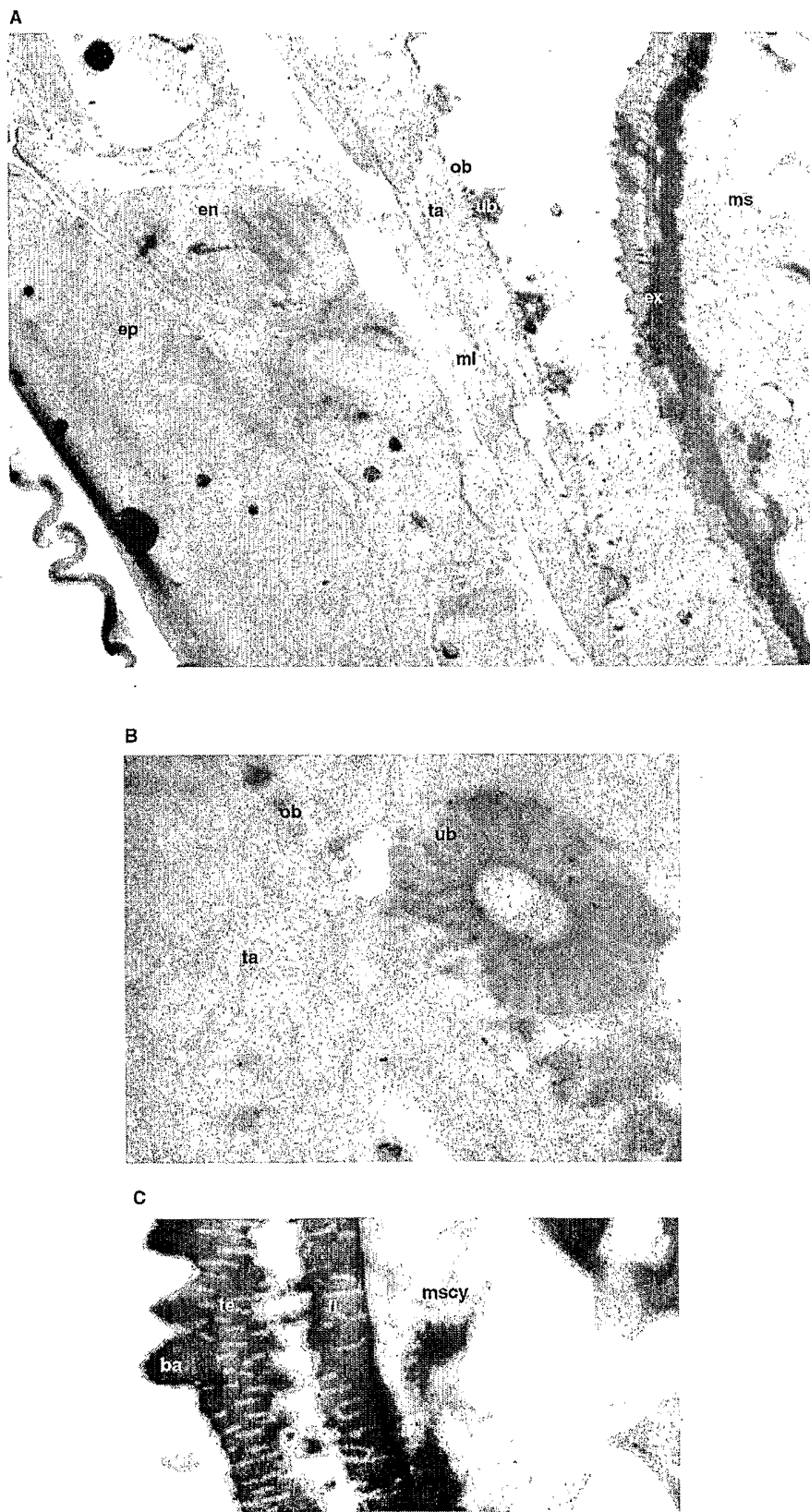
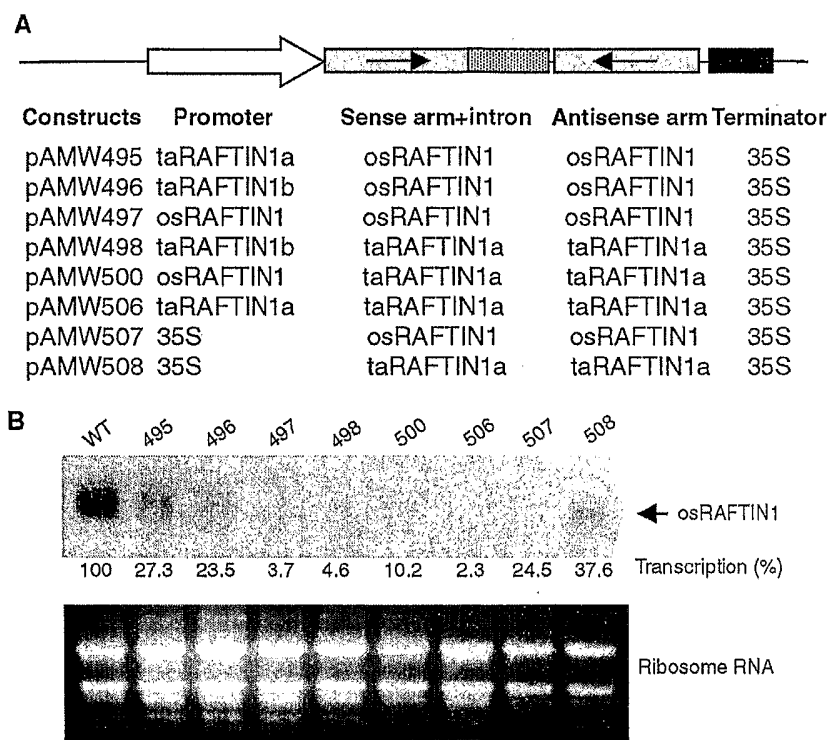


Figure 6

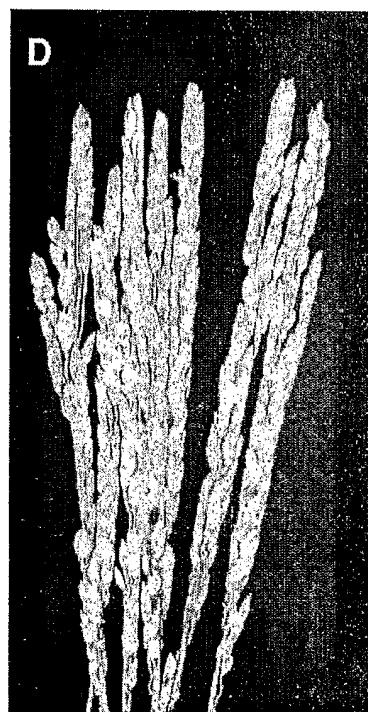
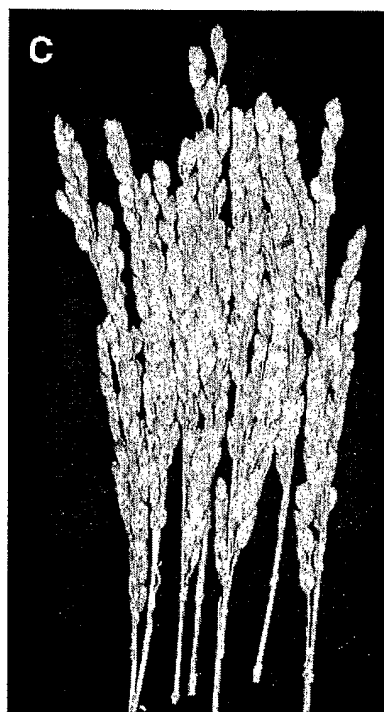
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Figure 7 Aiming Wang *et al.* 2002

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Figure 8ABCD Aiming Wang *et al.* 2002

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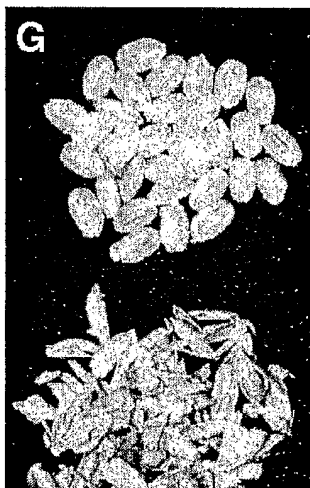
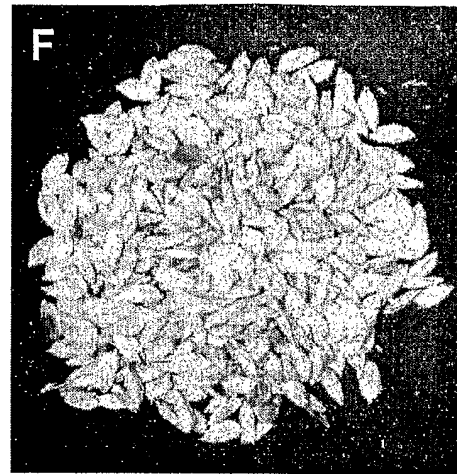
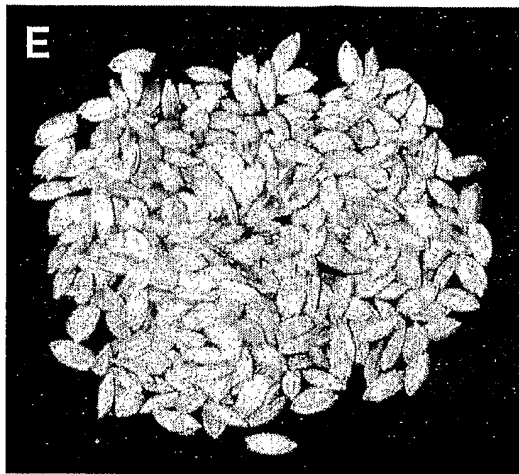


Figure 8EFGH Aiming Wang *et al.* 2002

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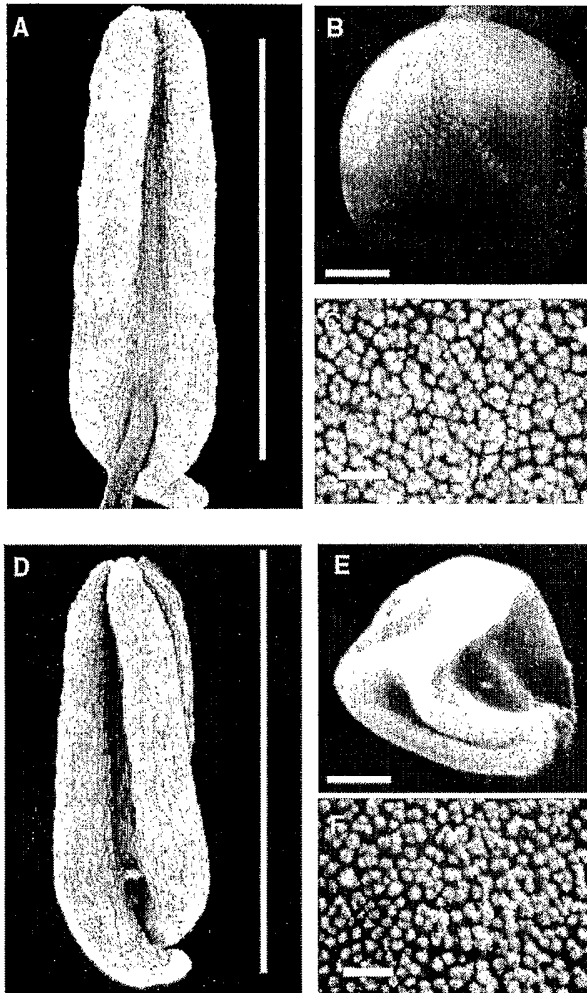
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Figure 8I Aiming Wang *et al.* 2002



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Top
↑

Figure 9 Aiming Wang *et al.* 2002

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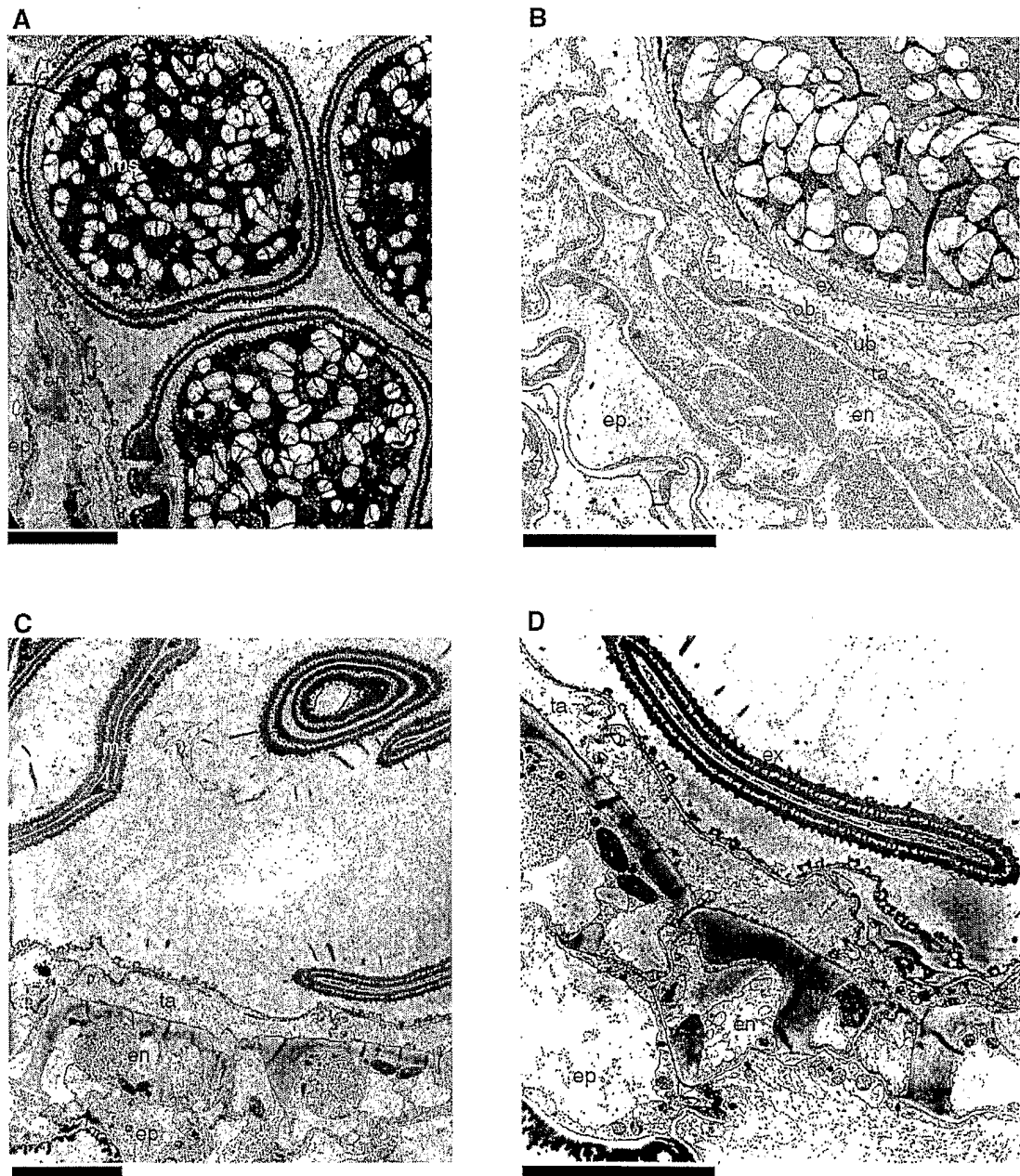


Figure 10

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Fig. 11. *taRAFTIN1a* cDNA sequence (1338 nt excluding the polyA tail, ORF from nt 29 to nt 1198). Start codon and stop codon are underlined.

CTCTGGACCTCTCACCTAGCGCACATCCATGGCGCGCTTCCTCGTCGCCCTCCTCGCCACCAC
CCTGGTCGCGGTTTCAGGCTGGAGGGCAGCTGGGGCCACGCGGCGCCGGCGACGGCGGAGGTGTT
CTGGCGCGCCGTGCTGCCCACTCGCCATTGCCCGACGCCGTTCTCCGCCTTCTCAAACAACC
CGCAGCAGGTGTTGAACTGCTCACAGAAGCCACCAGCTTCGTGAGGGATGCCGAGGACAGGCC
CCCCTTCGACTACCGTGATTACAGCCGCTCGCCGCCCGATGATGAACCGAGCAAGAGCACCGG
CGCCGCCTCCGGGGCGCGGGACTTCGACTACGACGACTACAGCGGGGGCGACAAGCTCCGTGG
CGCCGCCTCCGGGGCGCGGGACTTCGACTACGACGACTACAGCGGGGGCCGACAAGCTCCGTGG
CGCCACCGATGAATACAAGGCGCCGAGCAGCAGCCTCGCTGGAAACGGGGCGTCCATGGCTAG
GGGCGGCAAGGCGGAGACGACGACGGTGTCTTTCACGAGGAGGCGGTGCGCGTCGGCAAGAG
GCTCCCATTCGCTTCCCGCCGGCGACTCCCGCCGCGCTCGGTTTCCTGCCGCGCCAGGTGCG
CGACTCCGTCCCGTTACGACGGCCGCGCTGCCTGGCGTCCTCGCGACGTTCCGGCGTCGCGTC
CGACTCCGCCACGGTGGCCAGCATGGAGGCGACGCTGCGCGCCTGCGAGTCGCCGACCATCGC
CGGGGAGTCCAAGTTCTGCGCGACCTCGCTGGAGGCCCTGGTGGAGCGCGCCATGGAAGTGCT
GGGGACCCGCGACATCAGGCCGGTGACGTCGACGCTGCCCCGCGCCGGCGCCCCGCTGCAGAC
GTACACCGTCCGCTCCGTGCGGCCGGTGGAGGGGGGGCCTGTCTTCGTGGCGTGCCACGACGA
GGCCTACCCGTACACCGTGTACCGGTGCCACACCACTGGCCCGTCCAGGGCGTACATGGTGGA
CATGGAGGGCGCGCGCGGGCGGACGCGGTGACCATCGCCACCGTGTGCCACACCGACACGTC
CCTGTGGAACCCGGAGCACGTCTCCTTCAAGCTCCTGGGCACCAAGCCTGGCGGCACGCCGGT
CTGCCACCTCATGCCGTACGGGCACATAATCTGGGCCAAGAACGTGAATCGCTCGCCGGCGTG
AGCGGCCCGGGCAGCTCTGTGGTCTCGCCGGACTAAGATCGATGTACTACTACTACTATCTG
TTTCCACCTACGTCTTCTGTTGTTTCAGACCACCAGATGGTCACCAGAGCAGCGCTTGTAATAA
AAGAACAGCTTCTGCAAAAAAAAAAAAAAAAAA

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Fig. 12. *taRAFTIN1a* genomic sequence (1560 bps including two introns). Introns are shown in lower case letters. Start codon and stop codon are underlined.

CTCTGGACCTCTCACCTAGCGCACATCCATGGCGCGCTTCCTCGTCGCCCTCCTCGCCACCAC
CCTGGTCGCGgtaatggccgaagaagccactgagcaacgcctgcatcttcttcatttcggcaa
actgcacctagtgcatttcgcatgagattgatcgatcacaaactgggtgctaacggcctgtttc
gtcacagGTTTCAGGCTGGAGGGCAGCTGGGCCACGCGGCGCCGGCGACGGCGGAGGTGTTCTG
GCGCGCCGTGCTGCCACACTCGCCATTGCCCGACGCCGTTCTCCGCCTTCTCAAACAACCCGC
AGCAGgtctgtctttcatgttcctttcctcgtcgccctccgttaactgtcttcttctctcgag
tttgattgaccgccaacacaaaaaatgcatgcacgcacagGTGTTGAACTGCTCACAGAAGC
CACCAGCTTCGTGAGGGATGCCGAGGACAGGCCCCCTTCGACTACCGTGATTACAGCCGCTC
GCCGCCCGATGATGAACCGAGCAAGAGCACCGGCGCCGCTCCGGGGCGCGGGACTTCGACTA
CGACGACTACAGCGGGGGCGACAAGCTCCGTGGCGCCACCGATGAATACAAGGCGCCGAGCAG
CAGCCTCGCTGGAAACGGGGCGTCCATGGCTAGGGGCGGCAAGGCGGAGACGACGACGGTGTT
CTTTACGAGGAGGCGGTGCGCGTCGGCAAGAGGCTCCCATTCGCTTCCCGCCGGCGACTCC
CGCCGCGCTCGGTTTCTGCGCGCCAGGTGCGCGACTCCGTCCCGTTACGACGCGCCGCGCT
GCCTGGCGTCTCTCGCGACGTTGCGCGTCGCGTCCGACTCCGCCACGGTGCCAGCATGGAGGC
GACGCTGCGCGCCTGCGAGTCGCCGACCATCGCCGGGGAGTCCAAGTTCTGCGCGACCTCGCT
GGAGGCCCTGGTGGAGCGCGCCATGGAAGTGCTGGGGACCCGCGACATCAGGCCGGTGACGTC
GACGCTGCCCCGCGCCGGCGCCCCGCTGCAGACGTACACCGTCCGCTCCGTGCGGCCGGTGGA
GGGGGGGCCTGTCTTCGTGGCGTGCCACGACGAGGCCTACCCGTACACCGTGTACCGGTGCCA
CACCCTGGCCCCGTCCAGGGCGTACATGGTGGACATGGAGGGCGCGCGCGCGGCGACGCGGT
GACCATCGCCACCGTGTGCCACACCGACACGTCCCTGTGGAACCCGGAGCACGTCTCCTTCAA
GCTCCTGGGCACCAAGCCTGGCGGCACGCCGGTCTGCCACCTCATGCCGTACGGGCACATAAT
CTGGGCCAAGAACGTGAATCGCTCGCCGGCGTGAGCGGCCCGGGCAGCTCTGTGGTCTCGCCG
GAACTAAGATCGATGTACTACTACTATCTGTTTCCACCTACGTCTTCTGTTGTTTCAGACC
ACCAGATGGTCACCAGAGCAGCGCTTGTAAATAAAAGAACAGCTTCTGC

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Fig.13. *taRAFTIN1a* promoter sequence (1719 bps).

CTGTGATGGCGCTCTGTCCTTGTGATTCTTTCTTAGGGAACTCGTCTCTGGGGCCTCCGAGG
CCTGCAACCCTGTATCAGGACAAATTCTGACTGGCCTCCAGGAGTCCTAACAGCCACCGACCTG
GTCCACTGGGCCCCTAGAGTATCTTGAAGTGTCGTTTGCACAAATCCCGCTAATTAAGGGA
TGTGATGATGATGGTTTCTGAATCCGCGCGCCTTACCTCGCAAAACGGGGAATTGCAAAGGAT
ATATGGCACCTGTCGCGTCGTGAGGCCAGACGCTTCGGTTTCAAGCTGGTTATAGGGAGGGGG
AAACGAAGGGTTTTTTCTCCCTCTGTCTTCATCCATTTTCGTCTCCCAGCCCTCAGCTCCCAA
AAGCGTGTCGCCACCTCAAAGTCTTCAGCGCTTGCTCACGTAGCCCCCGTCCACCCCTTCCTT
GCCACCAAGATGGCCCGAACCAAGAGCGAGAAGGTTCCCTAAGGTTCCCAGCTAGGATCTGCCC
GCCGCTGGAACGGGGCTGAAGCGGAAGAGGGTTCGCTCCAAGGGTGGTATGAAACAACAGCCG
GAAGCCCCCAAGACTACAGGAAAGTGGTTCCTTCCTCGGCCACCGACAAAAAATTTCAGGGT
CTCGTGAGATAGGGCTGATGCCAGCGGATTTGGAGTGCCGCTCCCGGGGGACGAGGCTCCG
CCAACTCCTCGCGACGGTGAGCACATCCTCTGCCTGGAGTATATATTTTCGGGAGGGGGCTCGGG
TTTCCCCTACACGACTTCGTTTGCGGGATCTTGCGCTTCTACGGCTGCTAGCTACACCACATC
CCGTCAAACGGGGTTCTTTACATTGCAAACCTTCATCACATTTTGCGAGTGCTTTCTCGGGACT
GCCGCTCACTTTAAGTTGTTCCAATACTTCAATCAGGACTGCGTTCAGACCAACGGGGACATC
GTCTACGACCCCGCAACACCAAAATTCCTCGCCACATACCTCCGAAAATAATCCTATACAACC
TGGTCTCACGCTTCATCTCGTAAGATTTGCCATGTGTACTTCACCAATCTTGATGCATCCCTT
TTTCCCCAAGATTTATATGCCTGATCTGTATTTTGTCTCCGCTGTTTTCGAGATTTGATGTTTA
ATTGATGAAGCCCAAGCAATCCGGCATGCCCGTCGGTGCACTAGATGGCTAGCTTTTCTACGG
TGCTGGGCCTGCCGGCGAGGGGCGCGAGGCCACGTAGGAGACTGTTAGGATTCATGGGGCTGG
ACGCTGGTGGCGTGAAGTTTCGGGAAGGAGGATTGAGGAAGAAGGATGCATCAAGATTGGTGAA
GAACACGTGGCATCCTCTAGAGTAGGTCTTACGAGATGAAGCCTGAGACCAGGTCGTATGGGA
TTATTTTCCCGGACCTCCCGAAGCCCCGCAAAGTTAACTGCAGCTGCGTGGACGGCGAGCACC
GCACCGCACACGAACGCGAACCTGACGCTGCCGCGCCACACAACACGCCATTTCGCGCGCGGAT
CGTCGGATGTCACGCCCAGGATTATATTTCTCCGGTGCCGCACGTACCATGCGATCGCACAGCT
CACGTCGAGAGCTTTTCTGTTTGGCGTCGCGTCAATGAAACACCTTCCCGTCGAGCCGACGA
CGCCTATAAGTACCTCGTCTGATCGCATCATCACTCCCAAGTACTACAACCTCTGGACCTCTC
ACCTAGCGCACATCCATG

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Fig.14. *taRAFTIN1b* cDNA sequence (1275 bps excluding the polyA tail, ORF from nt 25 to nt 1113). Start codon and stop codon are underlined.

CGACCTCTCACCTAGCGCACATCCATGGCGCGCTTCCTCGTCGCCCTCCTCGCTGCCACCCTG
GTCGCGGTTTCAGGCTGGAGGGCAGCTGGGCCACGCGGCGCCGGCTACGGGGGAGGTGTTCTGG
CGCGCCGTGCTGCCGCACTCGCCATTGCCTGACGCCGTTCTCCGCCTCCTCAAACAACCTGCA
GCAGAATCCACCAGCTTCGTGAGAGACCCCGAGGACAGGCCCCCTTCGACTACCGTGATTAC
AGCCGCTCGTCGTCCGATGATGAACCGAGCAAGAGCACCGTCGCCGCTCCGGAGCGGGGGGC
TTCGACTACGACAACTACAGCGGGGCCGACGAACGTCGTGGTGCCACCGATGAATACAAGGCG
CCGAGCAGCAGCCTCGCTGGAAGCGGGGCGTACATGGCTAGGGGCGGCAAGGCGGAGACGACG
ACGGTGTTCTTTCACGAGGAGGCGGTGCGCGTCGGCAGGAGGCTCCCATTCCACTTCCCGCCG
GCGACTCCCGCCGCTCTCGGTTTCCTGCCGCGCCAGGTGCGCGACTCCGTCCCGTTACGACG
GCCGCGCTGCCCGGCATCCTCGCGACGTTTGGCATCGCGTCCGACTCCACCACGGTGCCGAGC
ATGGAGGCGACGCTGCGCGCCTGCGAGTCGCCCACCATCGCCGGGGAGTCCAAGTTCTGCGCG
ACTTCGCTGGAGGCCCTGGTGAGCGCGCCATGGGAGTGCTGGGGACCCGGGACATCAGGCCG
GTGACGTCGACGCTGCCCCGCGCCGGCGCCCCGCTGCAGACGTACACCGTCGTCGCCGTGCAG
CCGGTGGAGGGGGGGCCTGTCTTCGTGGCGTGCCACGACGAGGCCTACCCGTACACCGTGTAC
CGGTGCCACACCACCGGCCCGTCCAGGGCGTACACGGTGGACATGGAGGGCGCGCGCGCGGCC
GACGCGGTGACCATCGCCGCCGTGTGCCACACCGACACGTCCCTGTGGAACCCGAGCACGTC
TCCTTCAAGCTCCTCGGCACCAAGCCCGGCGGCACGCCGGTCTGCCACCTCATGCCGTACGGG
CACATAATCTGGGCCAAGAACGTGAAGCGCTCGCCGGCGTGAGCGGCCTTGACGCTCTGTGGT
GTCGCCGGAATAAGATCGATGTACTACTACTATCTGTTCTTACCTACGTCTTCTTGTTGTTT
ATACCACCAGATGGTCACCCAAGAGCAAGCGTTCGTAATAAAAAGAACAGCTTTTTTGCAGAAG
CTGGTGTTTTATTTTAAAAA

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Fig.15. *taRAFTIN1b* genomic sequence (1503 bps including two introns). Introns are shown in lower case letters. Start codon and stop codon are underlined.

CGACCTCTCACCTAGCGCACATCCATGGCGCGCTTCCTCGTCGCCCTCCTCGCTGCCACCCTG
 GTCGCGgtaatggccgaagaagagcaacgcctgcatcttcttcattttggcaaattgcaccta
 gtacattttgcatgagattaatcaatcacaaactggtgctaacggcctgtttcgtcccag
 GTTCAGGCTGGAGGGCAGCTGGGCCACGCGGCGCCGGCTACGGGGGAGGTGTTCTGGCGCGCC
 GTGCTGCCGCACTCGCCATTGCCTGACGCCGTTCTCCGCCTCCTCAAACAACCTGCAGCAGgt
 ctgtcttgcatcttctcgtcgccttcggttaactgtcttcttctctcgagtttgattgatca
 ccaaacacaaaaatgcatgcacgcgtgggtgttgaaactgcgcacagAATCCACCAGCTTCGTG
 AGAGACCCCGAGGACAGGCCCCCTTCGACTACCGTGATTACAGCCGCTCGTCGTCCGATGAT
 GAACCGAGCAAGAGCACCGTCGCCGCCTCCGGAGCGGGGGGCTTCGACTACGACAACCTACAGC
 GGGGCCGACGAACGTCGTGGTGCCACCGATGAATACAAGGCGCCGAGCAGCAGCCTCGCTGGA
 AGCGGGGCGTACATGGCTAGGGGCGGCAAGGCGGAGACGACGACGCGTGTTCCTTTCACGAGGAG
 GCGGTGCGCGTCGCGCAGGAGGCTCCCATTCACCTTCGCGCGGCGACTCCCGCCGCTCTCGGT
 TTCCTGCCGCGCCAGGTGCGCCGACTCCGTCCCGTTCACGACGGCCGCGCTGCCCGGCATCCTC
 GCGACGTTTGGCATCGCGTCCGACTCCACCACGGTGCCAGCATGGAGGCGACGCTGCGCGCC
 TGCAGATCGCCACCATCGCCGGGAGTCCAAGTTCTGCGCGACTTCGCTGGAGGCCCTGGTG
 GAGCGCGCCATGGGAGTGCTGGGGACCCGGGACATCAGGCCGGTGACGTCGACGCTGCCCCGC
 GCCGGCGCCCCGCTGCAGACGTACACCGTCGTGCGCGTGCAGCCGGTGGAGGGGGGGCCTGTC
 TTCGTGGCGTGCCACGACGAGGCCTACCCGTACACCGTGTACCGGTGCCACACCACCGGCCCG
 TCCAGGGCGTACACGGTGGACATGGAGGGCGCGCGCGGCGCCGACGCGGTGACCATCGCCGCC
 GTGTGCCACACCGACACGTCCCTGTGGAACCCGGAGCACGTCTCCTTCAAGCTCCTCGGCACC
 AAGCCCGGCGGCACGCCGGTCTGCCACCTCATGCCGTACGGGCACATAATCTGGGCCAAGAAC
 GTGAAGCGCTCGCCGGCGTGAGCGGCCTTGACGCTCTGTGGTGTGCGCCGGAACCTAAGATCGAT
 GTACTACTACTATCTGTTCTTACCTACGTCTTCTTGTGTTCATACCACCAGATGGTCACCCA
 AGAGCAAGCGTTCGTAATAAAAAGAACAGCTTTTTTGCAGAAGCTGGTGTTTTATTTT

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Fig. 16. *taRAFTIN1b* promoter sequence (2095 bps).

TTGTTGAGTGCCACACTATATTTCACTACACCATATGCACATTATGCTTGGATTGTCTTGTACT
TGACTCATGTGTTTAGACACTTCATTTTATTTGGTGTGTGAATGACTCCTATGCTTACCATA
GACCTTTCATTGAGCGCTTTGTGCATGTTTGTATACCTTGAGGTAGATGTTTGTCTCTTGT
CAAATATATAGCATCTCTACCTCCCATTTGCATGCTTGTTCATGATGTCCTTGATTGTGCT
CAATTCATATGCTTCTGTGACATGCCACAATCCTTTGTACACCATATGCTAGGCTTGATGAT
GACACTTGTGTTGGGTGACTCACCTTTTGAATGATTGGTTTTGCATTAACGCTAACCACATTTAT
TTTTCCAAGTGTTTGTGTCCTTGCTCCTTTTGAAGGAACCACATGACGGTGCGACATTGGAG
AGTGCCTATTTTCGAGCTTCAAGATGATGAGTGCTTGGTGATCGTCCACTTCTACATGGTGACG
CCGTCTCTTTCCCATGGTGATTGTTGGTTTTTGATCCGAGGTCGGATCTTTCCCAAGTGGGAGGG
GATGATGCGGAGCATACTACGGACATCACCATGTCTAGAGTTCATTACAGCAAGTGACACCTAT
CACATCTACTTCACATACATAAAGGTGAATCATCTCCTTTACACGTGCTCACTTGATCCCTTC
GAGGATGGTATACTACTTGACACTTCTCACGTGTGCATGCATAGGCATTGTGCGGAGCACCATG
AACGATGAGGAGGAGTGCGGAGCACAAGTGTACAACACACCATCCGCGAGGGAAGCATGGAAG
AGAAGGAAGAAGAAGCATGGACAAGCTTCTGGAAAGCCCGGAACCTTCTGGCCTCCTGCCCGGA
ACTTCCGGTCATCCGAAACTTCTGCCCCGACACACCGAAGCCGTCTGAGAGCGTGCCAAATC
TCTGGATAGCCCGGACCTTCGACCGGAACCTCCGGCGCCTGGACCTTCCGGCCATCCCTGGAA
CTCCCGGCCTGCCTGCACGCAGAGACTCGGGCCGAAGCGCATGTACCCTTTGCCCCCTCACTT
ATCCCTTCGTGGCTATCACTATATATACTCATCCTCCTCCTCCATTCTAGGGTTAGCATTTTG
ATAGCTCATTTGCATGTGAGATTTGCTCCTTACCCCCATCTCCTCTTGAGAGAGTGAGATTGA
TGCACTCCATTGGAGTCCAAGGTCTCCTTTGGAGAAGATCCCATAGGGGAATCAAGACCCCAT
CATGGGAAGATCCTTCTAGGATTCAAGACCTCAACTCCTTTAAGGATTGGGATGAACTAGTTA
CCTCTTGATCTTCTTGTGTTGGATTTAAACCTTTGTATCCCTCTATGTGTATGTGGATTTAG
CATATGTGTGATTGGATCTTGTCTATTGGAGTGTTTCTCTCTTTTGTTCCTTGTGTTTCAT
CGTTTTCTTCGGGAGATCCCCTCCATTTCTGTGAAAGATCGGTCCCTAGGGTTCTACCCCTACAT
TAGCTCAGGTTTTCCCTACACATCTTCGTTTGTGAGCTGTTGCGCTTCTACGGCTGGGAGCTA
CAGCACATCTCATTTCCACCAACGGGGTTCTTCACATTGTAAACTTCATCGTATTTTGCGAA
TGCTTTCTGGGGACAGCCACTCACTTTGAGTTGTTCCGATACTTCTTCCGGGTCTGCGTTTCAG
ACCAACGGGGACACCGTCTGCAACCTTGGAGGAGCCATTCTGCGACACACCAAAATTTTCGC
CACGGACCCCCCGAAGATCCGCAAGAAAAAAAAGCTGCAACGGCGTGAGCGGCGAGCACCGC
ACCGCACACGAACGCGAACGCGACGCTGCCGCGCCACACAACACGCCATTGCGCGCGCGGATCG
TCGGATGTCACGCCCACGATAATATTTCTCCGGTGCCGCACGTACCATGCGATCGCACAGCTCA
CATCGAGAGCTTTTCTGTTTGGTGTGCGCGTCAATGAAACACCTTCCCGTCAAGCCGACGACG
CCTATAAGTACCTCGCCTGATCGCATTATCACTCCCAAGTACTACAACCTCTCGACCTCTCAC
CTAGCGCACATCCATG

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Fig. 17. *taRAFTIN1d* predicted cDNA sequence (246 bps).

ATGGCGCGCTTCCTCGTCGCCCTCCTCGCTGCCACCCTGGTCGCGGTTTCAGGCTGGAGGGCAG
CTGGGCCACGCAGCGCCGGCGACGGCGGAGGTGTTCTGGCGCGCCGTGCTGCCGCACTCGCCA
TTGCCCCGACGCCGTTCTCCGCCTCCTCAAACAACCTGCAGCAGGTGTTGAACTGCACACAGAA
GCCACCAGCTTCGTAAGAGACCCCGAGGACAGGCCCCCCTTCGACTACCGTGATTAC

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Fig. 18. *taRAFTIN1d* partial genomic sequence (441 bps). Introns are shown in lower case letters.

ATGGCGCGCTTCCTCGTCGCCCTCCTCGCTGCCACCCTGGTCGCGgtaatggccgaagaagcc
actgagcaacgcctgcatcttctttatTTTggcaaactgggtgctaacggccaatactgccgct
tgcgttacgtctcagGTTCAAGCTGGAGGGCAGCTGGGCCACGCAGCGCCGGCGACGGCGGAG
GTGTTCTGGCGCGCCGTGCTGCCGCACTCGCCATTGCCCGACGCCGTTCTCCGCCTCCTCAAA
CAACCTGCAGCAGgtctgtcttgcattgctcctcgtcgccctccgttaactgtcttcttctctc
gagtttgattgatcaccaaacacaaaaatgcatgcacgcgtacgcgtagGTGTTGAACTGCAC
ACAGAAGCCACCAGCTTCGTAAGAGACCCCGAGGACAGGCCCCCTTCGACTACCGTGATTAC

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Fig. 19. *osRAFTINI* cDNA (1301 bps, ORF from nt 63 to nt1301). Start and stop codons are underlined.

GTTCGCAGTCGTCTCCGGCGAGAAATCGGCTGCGCCCCGTCTCTCTCTCTCTCGAACGCTTCCA
FGGCGCGCTTCCTCCTCCTCCTCGTCGCCGTGCGCCGTGCGCCGCCGTGCTTTCGCTGGGCG
ACGCGGCGCCGTGACGCGCCGAGGTGTTCTGGCGCGCCGTGCTGCCGGAATCCCCGTTGCCGG
ACGCCTTCCTCCGCCTCCTCCGCCCTGACACCAGCTTCGTTCGTTCGGCAAAGCGGAGGCGGCCG
GTGGCGCGGCGCGGACCGGATTCCCCTTCGATTACACTGACTACAGGGGATCTGATTCTCCGA
CGACGGCGAGTGTTTGGACCTCGCCCGTGACTTCGGCGAGCCGGCGCCTTTTCGGCTACGACT
ACAGTGACAGGGCGAAGGCGGCGGCGGCGGCCGCCGCCGCCGCCGGAGAGCAGGTTCTTG
CCGTCGACGCGGGCTTCAACTACGACAAATACGTTCGGCGCGAGGAAGCTCCGCGGCGGCAGCA
GCACCGCCGGCGGAGAGAATGATGACGAGCCTTTCGGGTACGACTACAAGGCGCCGAGCAGCG
GCAGCGGCACCGCGGCGTCGACGACGGCGCGAGGCGTCGGCACGGGCGCCACGACGACGGTGT
TCTTCCACGAGGAGGCGGTGCGCGTCGGCGAGAGGCTCCCGTTCTACTTCCCGGCGGCGACGA
CGTCGGCGCTGGGCTTCCTGCCGCGCCGCGTCGCGGACTCCATCCCGTTCACGGCGGCGCGCG
TGCCGGCCGTCTTCGCGCTGTTTCGGCGTCGCGCCGGACACCGCCGAGGCGGCGGCATGAGGG
AGACGCTGCGCACGTGCGAGTGCGCGACCCCTCGCCGGCGAGTCCAAGTTCTGCGCCACGTTCGC
TGGAGGCCCTGGTGGAGGGCGCCATGGCGGCGCTCGGGACACGCGACATCGCCGCGCTGGCGT
CGACGCTGCCCCGCGGCGGCGCGCCGCTGCAGGCGTACGCCGTCCGCGCCGTGCTCCCCGTG
AGGGCGCCGGCTTCGTGGCGTGCCACGACCAGGCGTACCCGTACACCGTGTACCGCTGCCACA
CCACCGGCCCGGCCAGAGCTTACATGGTGGAGATGGAAGGCGACGGCGGCGGCGATGGCGGCG
AGGCGGTGACCGTGGCCACCGTGTGCCACACCAACACGTTCGCGGTGGAACCCGGAGCACGTCT
CGTTCAAGCTCCTCGGCACCAAGCCCGGCGGCTCGCCGGTGTGCCACCTCATGCCGTACGGGC
ACATCGTCTGGGCCAAGAACGTGAAGAGCTCGACGGCGTAG

[illegible]

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Fig. 21. *osRAFTIN1* promoter sequence (1461 bps).

CGAAGGCAAACCTCTGGTAAGGATTCCCATTTACACGAATCAATTTAATAAGTCTAAAACGAACA
CTATGTTATGAGAAACACCTCACATCCGTCCATAACCGTGGGCATGACTATTTAAAAAGTTTA
ACTAAACTCTACAAAAGTTGCACGCTTTACCCACACGTCATGAACGTTTCACATTACCGAATA
CATGTGGATCGGACATGGCCGACAAAGGAGAGTTCAATACAAGGCTTTTCCATAACCAATCCA
TAAATATCCTATGTCCCACGGTTGGGTGGAATCTCTCCACCAAACATCAAGCCAGGATCAGGT
CCTCATCTACCCATGCCCCACTCCATGGACTCCGACACATCCCCACTGCAGGAGATTGCCATA
TACGCCACCATAACAGTGCTCCTCAACCGCTAACATGTTGGACACCAAATTCTATATACTTAT
ATAGTTCATCTCCACTAAGTGTAGTTAATTACATTTCTCTCTTCTCTCATTAAGCCACATCAC
CTCAATTATTTTTCAGCCTTTAGATGATAGATCTATGGTCCAAATTGTCTTTTCTTCTCTCT
CTTAAAAACATGCAATCTTAAATACTTTTAGGCTCAAATTTGTATCAAATTGTTTTAGTTTTG
TACATATTATGCAACTTAATTTTTCGCCGCAACGCGGAGGGGTATTTTCATCTAGTATTATTTA
AGAGCTATACACACTGCTATAGGGGAAAAAAAAGATAGGTTTGGCCCCCTGGTCAGTCCTGTT
GCACGGCTATATGTTGAAGGGAAAAAGCCAGTACGTTTTGTAGGTTGTTTTTTTTTTAGAAAT
GCTAAAAAGTTGTGGCATGTTTTTTTAGGTAAAAGCCTTTAAATATAAGTTACATTGTAACAC
AGTGTAATTCGCTGTAACATATATTGTAATCTCTATATAAGTTAGATATAAAATTACATATAT
ATTATTTTAATACTTATTTATAAGTTAGTATATTATAGTTATAATGGAATTAATTATAATTAT
AGTATAGTTAGATTTGAAAGTTTTTCCTTTAAGAAATTTTCGCAACAGTTTATTAGATATAGTC
CCTAAACGAAAATGTCAGGTGGATGCATGATTTCAGTGTGACGCTCGGGCGGATCACGGCTGCG
TCACGAAAATTCCCCCATGCAACCCGCGTCCGGCCGCTCCTTCGTGCCAACAGGCAACAGCGC
GGCGCCGGCGAACGTCACGCCCAAGATTATATTCCCCCTCTCGCGCTCGCGCGCGCCGCGACG
TCGTCCGAGCCAACATTATTTTTCTGTTTCCTGTACCCGTCGCCGTTGATCTCAAGCGAGATT
TGAGGTTTGGCCACGACGACGCTGCCTATAAAATACCAGGTGGTGGTCACCGCCCGCGCGCGT
CGATCGATCCGTCGCAGTCGTCTCCGGCGAGAAATCGGCTGCGCCCCGTCTCTCTCTCTCG
AACGCTTCCATG

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Fig. 22. Predicted protein sequences

taRAFTIN1a (389 residues)

MARFLVALLATTLVAVQAGGQLGHAAPATAEVFWRAVLPHSPLPDAVLRLLKQPAAGVELLTEATSFVR
DAEDRPPFDYRDYSRSPDDEPSKSTGAASGARDFDYDDYSGGDKLRGAASGARDFDYDDYSGADKLRG
ATDEYKAPSSSLAGNGASMARGGKAETTTVFFHEEAVRVGKRLPFRFPPATPAALGFLPRQVADSVPFT
TAALPGVLATFGVASDSATVASMEATLRACESPTIAGESKFCATSLEALVERAMEVLGTRDIRPVTSTL
PRAGAPLQTYTVRSVRPVEGGPVFVACHDEAYPYTVYRCHTTGPSRAYMVDMEGARGGDAVTIATVCHT
DTSLWNPEHVSFKLLGTKPGGTPVCHLMPYGHIIWAKNVNRSPA

taRAFTIN1b (362 residues)

MARFLVALLAATLVAVQAGGQLGHAAPATGEVFWRAVLPHSPLPDAVLRLLKQPAAESTSFVRDPEDRP
PFDYRDYSRSSDDEPSKSTVAASGAGGFDYDNYSGADERRGATDEYKAPSSSLAGSGAYMARGGKAET
TTVFFHEEAVRVGRRLPFHFPPATPAALGFLPRQVADSVPFTTAALPGILATFGIASDSTTVPSMEATL
RACESPTIAGESKFCATSLEALVERAMGVGLTRDIRPVTSTLPRAGAPLQTYTVAVQPVVEGGPVFVAC
HDEAYPYTVYRCHTTGPSRAYTVDMEGARGADAVTIAAVCHTDTSLWNPEHVSFKLLGTKPGGTPVCHL
MPYGHIIWAKNVKRSPA

taRAFTIN1d (partial sequence, 82 residues)

MARFLVALLAATLVAVQAGGQLGHAAPATAEVFWRAVLPHSPLPDAVLRLLKQPAAGVELHTEATSFVR
DPEDRPPFDYRDY

osRAFTIN1 (412 residues)

MARFLLLLAVAAAAAVLSLGDAAPSTAEVFWRAVLPEPSPDAFLRLLRPDTSFVVGKAEAAAGGAART
GFPPFDYTDYRGSDSPTTASGLDLAGDFGEPAPFGYDYSAQGEGGGGAAAAAGEQVLAVDAGFNVDKYV
GARKLRGGSSTAGGENDEPFYDYKAPSSSGSGTAASTTARGVGTGATTTVFFHEEAVRVGERLPFYFP
AATTSALGFLPRRVADSIPTAAALPAVLALFGVAPDTAEAAAGMRETLRTCEWPTLAGESKFCATSLEA
LVEGAMAALGTRDIAALASTLPRGGAPLQAYAVRAVLPEVAGAGFVACHDQAYPYTVYRCHTTGPARAYM
VEMEGDGGGDGGEAVTVATVCHTNTSRWNPEHVSFKLLGTKPGGSPVCHLMPYGHIIWAKNVKSSTA

SEQUENCE LISTING

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<151> 2002-08-02

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17

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29

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ctccatgtcc accatgta 18

<210> 6
<211> 18
<212> DNA
<213> artificial

<220>
<223> Primer OL3148

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 <213> Triticum aestivum

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 <212> DNA
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<400> 31

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 <212> DNA
 <213> *Triticum aestivum*

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 <212> DNA
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<400> 33

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 <212> DNA
 <213> Triticum aestivum

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 <211> 246
 <212> DNA
 <213> Triticum aestivum

<400> 35

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 gattac 246

<210> 36
 <211> 441
 <212> DNA
 <213> Triticum aestivum

<400> 36

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<212> DNA

<213> *Oryza sativa*

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 ccgcgggaga gcaggttctt gccgtcgacg cgggcttcaa ctacgacaaa tacgtcggcg 480
 cgaggaagct ccgcggcggc agcagcaccg ccggcgggaga gaatgatgac gagcctttcg 540
 ggtacgacta caaggcgccg agcagcggca gcggcaccgc ggcgtcgacg acggcgcgag 600
 gcgtcggcac gggcgccacg acgacggtgt tcttccacga ggaggcgggtg cgcgtcggcg 660
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 gcgccatggc ggcgctcggg acacgcgaca tcgccgcgtt ggcgtcgacg ctgccccgcg 960
 gcggcgcgcc gctgcaggcg tacgccgtcc gcgccgtgct ccccgtcgag ggcgccgggt 1020
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 cggccagagc ttacatggtg gagatggaag gcgacggcgg cggcgatggc ggcgaggcgg 1140
 tgaccgtggc caccgtgtgc cacaccaaca cgtcgcggtg gaaccgggag cactctcgt 1200
 tcaagctcct cggcaccaag cccggcggct cgcgggtgtg ccacctcatg ccgtacgggc 1260
 acatcgtctg ggccaagaac gtgaagagct cgacggcgta g 1301

<210> 38
 <211> 1479
 <212> DNA
 <213> *Oryza sativa*

<400> 38

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gtcgcagtcg tctccggcga gaaatcggct gcgcccogtc tctctctctc tcgaacgctt      60
ccatggcgcg cttcctcctc ctctcgtcgc ccgtgcgcgc tgcgcgcgcg gtgcttttcgg      120
tacactcatg atgccgctac tcagctgagc catgcaccgt tgcacccgta tactaacgat      180
cgctcgatcg accgacgatg tgtgttcttc agcagctggg cgacgcggcg ccgtcgacgg      240
ccgaggtggt ctggcgcgcc gtgctgccgg aatccccgtt gcgggacgcc ttctccgcc      300
tctccgcccc tggtcgggtgt ccttccttcc tccttcgcgc gcgcgcgcgc gccattactc      360
tctcgcaggt ttgatttggt tgtggacgtt gcagacacca gcttcgtcgt cggcaaagcg      420
gaggcgggcg gtggcgcggc gcggaccgga ttccccttog attacactga ctacagggga      480
tctgattctc cgacgacggc gagtggtttg gacctgcgcg gtgacttcgg cgagccggcg      540
cctttcgggt acgactacag tgcacagggc gaaggcgggc gcggcgggcg cgcgcgcgc      600
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aggaagctcc gcggcggcag cagcaccgcc ggcgagaga atgatgacga gcctttcggg      720
tacgactaca aggcgccgag cagcggcagc ggcaccgcgc cgtcgacgac ggcgcgaggc      780
gtcggcacgg gcgccacgac gacggtgttc ttccacgagg aggcggtgcg cgtcggcgag      840
aggctcccg tctacttccc ggcggcgacg acgtcggcgc tgggcttccg gcgcgcgcgc      900
gtcgcggact ccatcccggt caccggcgcc gcgtgcgcgc ccgtcctcgc gctgttcggc      960
gtcgcgcgcg acaccgccga ggcgcccgcc atgagggaga cgtgcgcac gtgcgagtgg      1020
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gccatggcgg cgtcggggac acgcgacatc gccgcgctgg cgtcgacgct gcccgcgggc      1140
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gtggcggtgc acgaccaggc gtaccctgac accgtgtacc gctgccacac caccggcccg      1260
gccagagctt acatggtgga gatggaaggc gacggcggcg gcgatggcgg cgaggcggtg      1320
accgtggcca ccgtgtgcca caccaacacg tcgcggtgga acccgagca cgtctcgttc      1380
aagctcctcg gcaccaagcc cggcggtcgc ccggtgtgcc acctcatgcc gtacgggcac      1440
atcgtctggg ccaagaacgt gaagagctcg acggcgtag      1479

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<210> 39
 <211> 1461
 <212> DNA
 <213> *Oryza sativa*

<400> 39

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cgaaggcaaa ctctggtaag gattcccatt acacgaatca atttaataag tctaaaacga      60
acactatggt atgagaaaaca cctcacatcc gtccataacc gtgggcatga ctatttaaaa      120
agtttaacta aactctacaa aagttgcacg ctttaccac acgtcatgaa cgtttcacat      180
taccgaatac atgtggatcg gacatggccg acaaaggaga gttcaatata aggcttttcc      240
ataaccaatc cataaatatc ctatgtccca cggttgggtg gaatctctcc accaaacatc      300
aagccaggat caggtcctca tctacccatg cccaactcca tggactccga cacatcccca      360
ctgcaggaga ttgccatata cgccaccata ccagtgtctc tcaaccgcta acatgttgga      420
caccaaattc tatatactta tatagttcat ctccactaag tgtagttaat tacatttctc      480
tcttctctca ttaagccaca tcacctcaat tttttttagc ctttagatga tagatctatg      540
gtccaaattg tcttttcttt cttctctctt aaaaacatgc aatcttaa atcttttaggc      600
tcaaaattgt atcaaattgt tttagttttg tacatattat gcaactta attttcgccgc      660
aacgcggagg ggtatttcat ctagtattat ttaagagcta tacacactgc tataggggaa      720
aaaaaagata ggtttggccc cctggtcagt cctgttgca cggctatatgt tgaagggaaa      780
aagccagtac gttttgtagg ttgttttttt tttagaattg ctaaaaagtt gtggcatggt      840
ttttaggtaa aagccttta atataagtta cattgtaact acagtgtaat tccgctgtaa      900
ctatattgta atctctatat aagttagata taaaattaca tatatattat ttttaacttt      960
atttataagt tagtatatta tagttataat ggaattaatt ataattatag tatagttaga     1020
tttgaaagtt tttcctttaa gaaatttcgc aacagtttat tagatatagt ccctaaacga     1080
aaatgtcagg tggatgcatg attcagtgtg acgctcgggc ggatcacggc tgcgtcacga     1140
aaattccccc catgcaaccc gcgtccggcc gtcccttcgtg ccaacaggca acagcgcggc     1200
gccggcgaac gtcacgcca agattatatt cccctctctg cgctcgcgcg cgccgcgacg     1260
tcgtcggagc caacattatt tttctgtttc ctgtcaccgt cgccgttgat ctcaagcgag     1320
atttgagggt tggccacgac gacgcctgoc tataaatacc aggtgggtgg caccgcccgg     1380
cggcgtcgat cgatccgtcg cagtctctc cggcgagaaa tcggctgcgc cccgtctctc     1440
tctctctcga acgcttccat g                                     1461

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<210> 40

<211> 389
 <212> PRT
 <213> Triticum aestivum

<400> 40

Met Ala Arg Phe Leu Val Ala Leu Leu Ala Thr Thr Leu Val Ala Val
 1 5 10 15

Gln Ala Gly Gly Gln Leu Gly His Ala Ala Pro Ala Thr Ala Glu Val
 20 25 30

Phe Trp Arg Ala Val Leu Pro His Ser Pro Leu Pro Asp Ala Val Leu
 35 40 45

Arg Leu Leu Lys Gln Pro Ala Ala Gly Val Glu Leu Leu Thr Glu Ala
 50 55 60

Thr Ser Phe Val Arg Asp Ala Glu Asp Arg Pro Pro Phe Asp Tyr Arg
 65 70 75 80

Asp Tyr Ser Arg Ser Pro Pro Asp Asp Glu Pro Ser Lys Ser Thr Gly
 85 90 95

Ala Ala Ser Gly Ala Arg Asp Phe Asp Tyr Asp Asp Tyr Ser Gly Gly
 100 105 110

Asp Lys Leu Arg Gly Ala Ala Ser Gly Ala Arg Asp Phe Asp Tyr Asp
 115 120 125

Asp Tyr Ser Gly Ala Asp Lys Leu Arg Gly Ala Thr Asp Glu Tyr Lys
 130 135 140

Ala Pro Ser Ser Ser Leu Ala Gly Asn Gly Ala Ser Met Ala Arg Gly
 145 150 155 160

Gly Lys Ala Glu Thr Thr Thr Val Phe Phe His Glu Glu Ala Val Arg
 165 170 175

Val Gly Lys Arg Leu Pro Phe Arg Phe Pro Pro Ala Thr Pro Ala Ala
 180 185 190

Leu Gly Phe Leu Pro Arg Gln Val Ala Asp Ser Val Pro Phe Thr Thr
 195 200 205

Ala Ala Leu Pro Gly Val Leu Ala Thr Phe Gly Val Ala Ser Asp Ser

210 215 220
 Ala Thr Val Ala Ser Met Glu Ala Thr Leu Arg Ala Cys Glu Ser Pro
 225 230 235 240
 Thr Ile Ala Gly Glu Ser Lys Phe Cys Ala Thr Ser Leu Glu Ala Leu
 245 250 255
 Val Glu Arg Ala Met Glu Val Leu Gly Thr Arg Asp Ile Arg Pro Val
 260 265 270
 Thr Ser Thr Leu Pro Arg Ala Gly Ala Pro Leu Gln Thr Tyr Thr Val
 275 280 285
 Arg Ser Val Arg Pro Val Glu Gly Gly Pro Val Phe Val Ala Cys His
 290 295 300
 Asp Glu Ala Tyr Pro Tyr Thr Val Tyr Arg Cys His Thr Thr Gly Pro
 305 310 315 320
 Ser Arg Ala Tyr Met Val Asp Met Glu Gly Ala Arg Gly Gly Asp Ala
 325 330 335
 Val Thr Ile Ala Thr Val Cys His Thr Asp Thr Ser Leu Trp Asn Pro
 340 345 350
 Glu His Val Ser Phe Lys Leu Leu Gly Thr Lys Pro Gly Gly Thr Pro
 355 360 365
 Val Cys His Leu Met Pro Tyr Gly His Ile Ile Trp Ala Lys Asn Val
 370 375 380
 Asn Arg Ser Pro Ala
 385

<210> 41
 <211> 362
 <212> PRT
 <213> Triticum aestivum

<400> 41

Met Ala Arg Phe Leu Val Ala Leu Leu Ala Ala Thr Leu Val Ala Val
 1 5 10 15

Gln Ala Gly Gly Gln Leu Gly His Ala Ala Pro Ala Thr Gly Glu Val

Gln Thr Tyr Thr Val Val Ala Val Gln Pro Val Glu Gly Gly Pro Val
260 265 270

Phe Val Ala Cys His Asp Glu Ala Tyr Pro Tyr Thr Val Tyr Arg Cys
275 280 285

His Thr Thr Gly Pro Ser Arg Ala Tyr Thr Val Asp Met Glu Gly Ala
290 295 300

Arg Gly Ala Asp Ala Val Thr Ile Ala Ala Val Cys His Thr Asp Thr
305 310 315 320

Ser Leu Trp Asn Pro Glu His Val Ser Phe Lys Leu Leu Gly Thr Lys
325 330 335

Pro Gly Gly Thr Pro Val Cys His Leu Met Pro Tyr Gly His Ile Ile
340 345 350

Trp Ala Lys Asn Val Lys Arg Ser Pro Ala
355 360

<210> 42

<211> 82

<212> PRT

<213> Triticum aestivum

<400> 42

Met Ala Arg Phe Leu Val Ala Leu Leu Ala Ala Thr Leu Val Ala Val
1 5 10 15

Gln Ala Gly Gly Gln Leu Gly His Ala Ala Pro Ala Thr Ala Glu Val
20 25 30

Phe Trp Arg Ala Val Leu Pro His Ser Pro Leu Pro Asp Ala Val Leu
35 40 45

Arg Leu Leu Lys Gln Pro Ala Ala Gly Val Glu Leu His Thr Glu Ala
50 55 60

Thr Ser Phe Val Arg Asp Pro Glu Asp Arg Pro Pro Phe Asp Tyr Arg
65 70 75 80

Asp Tyr

<210> 43
 <211> 412
 <212> PRT
 <213> Oryza sativa

<400> 43

Met Ala Arg Phe Leu Leu Leu Leu Val Ala Val Ala Ala Ala Ala Ala
 1 5 10 15

Val Leu Ser Leu Gly Asp Ala Ala Pro Ser Thr Ala Glu Val Phe Trp
 20 25 30

Arg Ala Val Leu Pro Glu Ser Pro Leu Pro Asp Ala Phe Leu Arg Leu
 35 40 45

Leu Arg Pro Asp Thr Ser Phe Val Val Gly Lys Ala Glu Ala Ala Gly
 50 55 60

Gly Ala Ala Arg Thr Gly Phe Pro Phe Asp Tyr Thr Asp Tyr Arg Gly
 65 70 75 80

Ser Asp Ser Pro Thr Thr Ala Ser Gly Leu Asp Leu Ala Gly Asp Phe
 85 90 95

Gly Glu Pro Ala Pro Phe Gly Tyr Asp Tyr Ser Ala Gln Gly Glu Gly
 100 105 110

Gly Gly Gly Gly Ala Ala Ala Ala Ala Gly Glu Gln Val Leu Ala Val
 115 120 125

Asp Ala Gly Phe Asn Tyr Asp Lys Tyr Val Gly Ala Arg Lys Leu Arg
 130 135 140

Gly Gly Ser Ser Thr Ala Gly Gly Glu Asn Asp Asp Glu Pro Phe Gly
 145 150 155 160

Tyr Asp Tyr Lys Ala Pro Ser Ser Gly Ser Gly Thr Ala Ala Ser Thr
 165 170 175

Thr Ala Arg Gly Val Gly Thr Gly Ala Thr Thr Thr Val Phe Phe His
 180 185 190

Glu Glu Ala Val Arg Val Gly Glu Arg Leu Pro Phe Tyr Phe Pro Ala
 195 200 205

Ala Thr Thr Ser Ala Leu Gly Phe Leu Pro Arg Arg Val Ala Asp Ser
 210 215 220

Ile Pro Phe Thr Ala Ala Ala Leu Pro Ala Val Leu Ala Leu Phe Gly
 225 230 235 240

Val Ala Pro Asp Thr Ala Glu Ala Ala Gly Met Arg Glu Thr Leu Arg
 245 250 255

Thr Cys Glu Trp Pro Thr Leu Ala Gly Glu Ser Lys Phe Cys Ala Thr
 260 265 270

Ser Leu Glu Ala Leu Val Glu Gly Ala Met Ala Ala Leu Gly Thr Arg
 275 280 285

Asp Ile Ala Ala Leu Ala Ser Thr Leu Pro Arg Gly Gly Ala Pro Leu
 290 295 300

Gln Ala Tyr Ala Val Arg Ala Val Leu Pro Val Glu Gly Ala Gly Phe
 305 310 315 320

Val Ala Cys His Asp Gln Ala Tyr Pro Tyr Thr Val Tyr Arg Cys His
 325 330 335

Thr Thr Gly Pro Ala Arg Ala Tyr Met Val Glu Met Glu Gly Asp Gly
 340 345 350

Gly Gly Asp Gly Gly Glu Ala Val Thr Val Ala Thr Val Cys His Thr
 355 360 365

Asn Thr Ser Arg Trp Asn Pro Glu His Val Ser Phe Lys Leu Leu Gly
 370 375 380

Thr Lys Pro Gly Gly Ser Pro Val Cys His Leu Met Pro Tyr Gly His
 385 390 395 400

Ile Val Trp Ala Lys Asn Val Lys Ser Ser Thr Ala
 405 410