

Wuschel overexpression promotes somatic embryogenesis and induces organogenesis in cotton (*Gossypium hirsutum* L.) tissues cultured in vitro

O. Bouchabké-Coussa · M. Obellianne ·
D. Linderme · E. Montes · A. Maia-Grondard ·
F. Vilaine · C. Pannetier

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Abstract

Key message This work shows that overexpression of the *WUS* gene from *Arabidopsis* enhanced the expression of embryogenic competence and triggered organogenesis from some cells of the regenerated embryo-like structures.

Abstract *Agrobacterium*-mediated genetic transformation of cotton was described in the late 1980s, but is still time consuming and largely genotype dependant due to poor regeneration. To help solve this bottleneck, we over-expressed the *WUSCHEL* (*WUS*) gene, a homeobox transcription factor cloned in *Arabidopsis thaliana*, known to stimulate organogenesis and/or somatic embryogenesis in *Arabidopsis* tissues cultured in vitro. The *AtWUS* gene alone, and *AtWUS* gene fused to the *GFP* marker were compared to the *GFP* gene alone and to an empty construct used as a control. Somatic embryogenesis was improved in *WUS* expressed calli, as the percentage of explants giving rise to embryogenic tissues was significantly higher ($\times 3$) when *WUS* gene was over-expressed than in the control. An interesting result was that *WUS* embryogenic lines

evolved in green embryo-like structures giving rise to ectopic organogenesis never observed in any of our previous transformation experiments. Using our standard in vitro culture protocol, the overexpression of *AtWUS* in tissues of a recalcitrant variety did not result in the production of regenerated plants. This achievement will still require the optimization of other non-genetic factors, such as the balance of exogenous phytohormones. However, our results suggest that targeted expression of the *WUS* gene is a promising strategy to improve gene transfer in recalcitrant cotton cultivars.

Keywords Somatic embryogenesis · Organogenesis · *WUSCHEL* · *Gossypium hirsutum* · *Agrobacterium*-mediated transformation

Introduction

Transgenesis has been developed largely on the cotton plant, which is the most widely used source of vegetable textile fiber. Its economic importance is illustrated by the fact that 350 million people are employed in its production and manufacture. Given its economic importance and its vulnerability to insect pests, it is no surprise that cotton was one of the first crops studied with a view to conferring new agronomic traits through gene transfer. Genetically modified cotton is now widespread. Two-thirds (68 %) of the 36 million hectares under cotton cultivation were biotech in 2011 (source ISAAA: The International Service for the Acquisition of Agri-biotech Applications <http://www.isaaa.org>). Although results have been published on biolistic cotton tissue transformation (Finer and McMullen 1990; McCabe and Martinell 1993) and more recently by Liu and coll (Liu et al. 2011), until now the most commonly used

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O. Bouchabké-Coussa · M. Obellianne · D. Linderme ·
E. Montes · A. Maia-Grondard · F. Vilaine · C. Pannetier
INRA, UMR1318, Institut Jean-Pierre Bourgin, RD10,
78000 Versailles, France

O. Bouchabké-Coussa · M. Obellianne · D. Linderme ·
C. Pannetier
CIRAD, UR-SCA, 78026 Versailles, France

C. Pannetier (✉)
CIRAD UMR AGAP, 78026 Versailles, France
e-mail: catherine.pannetier@versailles.inra.fr;
catherine.pannetier@cirad.fr

process has been the *Agrobacterium*-mediated transformation and in vitro regeneration of transgenic plants. For cotton, the in vitro regeneration process is somatic embryogenesis which was first described in *Gossypium klotzschianum* (Price and Smith 1979). Following this pioneering work, several authors reported the regeneration of the cultivated cotton species *Gossypium hirsutum* through somatic embryogenesis (Trolinder and Goodin 1987; Gawel et al. 1986; Shoemaker et al. 1986). The Wilkins team developed highly regenerable elite Acala cotton (Mishra et al. 2003). This represents an important step towards genotype-independent regeneration, and hence, transformation. After the first reports of the production of transgenic cotton plants expressing the NPTII gene (Umbeck et al. 1987; Shoemaker et al. 1986; Firoozabady et al. 1987), the development of this technique to create new varieties expressing genes of interest (herbicidal or insect resistant) has been very rapid. *Agrobacterium*-mediated transformation of cotton and regeneration via somatic embryogenesis (SE) is being developed in several laboratories and private companies. Although several studies (Trolinder and Xhixian 1989; Cousins et al. 1991; Firoozabady and DeBoer 1993; Kumar et al. 1998; Sakhanokho et al. 2001, 2004; Sun et al. 2006; Zhang et al. 2009; Wu et al. 2004; Jin et al. 2006) reported the regeneration of various cultivars through somatic embryogenesis, many elite cultivars remain poorly regenerable (Mishra et al. 2003; Obembe et al. 2011). Following the first results published by Umbeck et al. (1987) and Firoozabady et al. (1987), the most efficient transformation process was developed for Coker varieties (Pannetier et al. 1997; Sunilkumar and Rathore 2001; Wilkins et al. 2004), and most transgenic cotton currently cultivated in the world came from primary transformants obtained on a Coker variety. However, the method is still time consuming mainly due to incomplete control of the regeneration process. Research into the regeneration process itself tends to be both tedious and protracted. For this reason as well as the economic premium attached to the rapid development of transgenic cotton, there have been few studies on the regeneration process and particularly on the induction of embryogenesis. The influence of media composition (mineral content and phytohormones) has been studied (Trolinder and Goodin 1988a, b). Transcriptomic approaches have been used to identify genes involved in somatic embryogenesis in cotton (Zeng et al. 2006; Wu et al. 2009; Yang et al. 2012) and more than 200 unigenes have been identified as upregulated during cotton SE (Zeng et al. 2006). It will be very challenging to identify among these genes those that regulate directly the process of somatic embryogenesis, and for evidence the recent work of Hu et al. (2011) is pioneer in this field.

Somatic embryogenesis is the biological process by which many plants can regenerate in vitro. In the past, most

studies focused on hormonal regulation of this process; (Zimmerman 1993; Lazzeri et al. 1987; Jimenez 2005) and the literature abounds with articles describing different strategies for regenerating a number of species via SE (Feher et al. 2003; Verdeil et al. 2007). More recently, due to progress made in studies of zygotic embryogenesis and of the shoot apical meristem a number of genes involved in SE induction have been identified (Verdeil et al. 2007; Tahir and Stasolla 2006; Rose and Nolan 2006). More specifically, many transcription factors: LEAFY COTYLEDON1 (LEC1) BABY BOOM (BBM) and AGAMOUS-LIKE 15 (AGL15) were shown to enhance embryo formation from vegetative cells, immature microspores or zygotic embryos (Alemanno et al. 2008; Lotan et al. 1998; Boutilier et al. 2002; Harding et al. 2003). We were interested in the effects of the *WUSCHEL* gene from *A. thaliana* expressed in cotton tissue cultured in vitro.

The *WUSCHEL* gene (*WUS*), which encodes a homeo-domain transcription factor, was initially identified as being required to maintain a pool of pluripotent stem cells in the shoot apical meristem (SAM) in an undifferentiated state (Endrizzi et al. 1996; Laux et al. 1996; Mayer et al. 1998). *WUS* expression is confined to a small group of cells in the lower part of the central zone of SAM, but it can drive signals across cell layers and is expressed non-autonomously (Mayer et al. 1998). *WUS* is thought to interact with *CLAVATA3*, a gene expressed in the underlying cell layers, by a regulatory loop controlling the size of the stem cell population, with the *CLV* genes repressing *WUS* at the transcript level, and *WUS* expression being sufficient to induce meristem cell identity (Brand et al. 2000; Schoof et al. 2000).

Ectopic expression of the *A. thaliana WUS* gene was shown to induce stem cells in vegetative tissues which can differentiate into somatic embryos without external plant hormones (Zuo et al. 2002). Other studies reported stem cell differentiation into organogenesis (Gallois et al. 2002). The ability of *WUS* to stimulate organogenesis and/or somatic embryogenesis appears to be dependant on the cellular context (Xu et al. 2005) or on the exogenous hormonal regime (Gallois et al. 2004). A connection was observed between *WUS* and cytokinins in the regulation of stem cells in the SAM (Leibfried et al. 2005). Proper apical meristem function requires the interaction between *WUS* and *Arabidopsis* response regulator (*ARR*) genes, which act in the negative feedback loop of cytokinin signaling, with *WUS* repressing the transcription of several *ARRs*. Thus, *WUSCHEL* facilitates high cytokinin activity (Shani et al. 2006). Cytokinin signals and *WUS* reinforce each other through multiple feedback loops (Sablowski 2009; Gordon et al. 2009). Su et al. (2009) showed a link between *WUS* and auxin. A correct *WUS* expression, regulated by a defined level of exogenous auxin concentration, is essential

for somatic embryo induction. Eventhough these studies concern de novo shoot formation, several papers highlight the role of *WUS* during in vitro regeneration (Cary et al. 2002; Gordon et al. 2007; Atta et al. 2009). All these results on the auxin–cytokinin–*WUS* connection, which clarify the role of hormones during in vitro shoot regeneration, along with previously published results on the ectopic expression of *WUSCHEL* leading to the formation of somatic embryos, provide supplementary arguments to perform experiments to analyze the effect of *WUS* over-expression on the cotton regeneration process through somatic embryogenesis.

Materials and methods

Plant material

Delinted cotton seeds of the Coker 310 variety were sterilized in a bayrochlor (Bayrol) solution (0.3 % active chlorine) for 30 min, rinsed with sterile water and sown in test tubes on a half strength MS medium (Murashige and Skoog 1962) supplemented with Morel and Wetmore vitamins (Morel and Wetmore 1951) 10 g/l sucrose and solidified with agar 8 g/L. Seedlings were grown in a culture room at $29^{\circ} \pm 1^{\circ} \text{C}$ under a low light intensity ($4 \mu\text{mol m}^{-2} \text{s}^{-1}$) for 5 days. Hypocotyl fragments of 4–5 mm in length were used as explants. Explants were placed on a basic medium (BM) composed of MS mineral salts (Murashige and Skoog 1962) supplemented with vitamins according to Morel and Wetmore (Morel and Wetmore 1951) and 20 g/L glucose, solidified with 4 g/L agarose (Litex, LSM 5000, Lonza Copenhagen). All the media used in our experiments had a pH adjusted at 5.8 and were autoclaved 20 min at 115°C . For callogenesis and embryogenesis induction, 2,4-dichlorophenoxyacetic acid (2,4-D) (0.1–0.05 mg/L) and kinetin (0.1–0.05 mg/L) were added. After 2–6 months, embryogenic clusters (Fig. 1c) appeared on the calli; they were excised and subcultured on a new medium to maintain embryogenic lines. The new medium was composed of the same Basic Medium containing 30 g/L sucrose instead of glucose and was free of phytohormones. Tissues were subcultured every 4 weeks. Control and *WUS*-overexpressing embryogenic clusters were isolated on the same basic medium.

Agrobacterium tumefaciens-mediated transformation

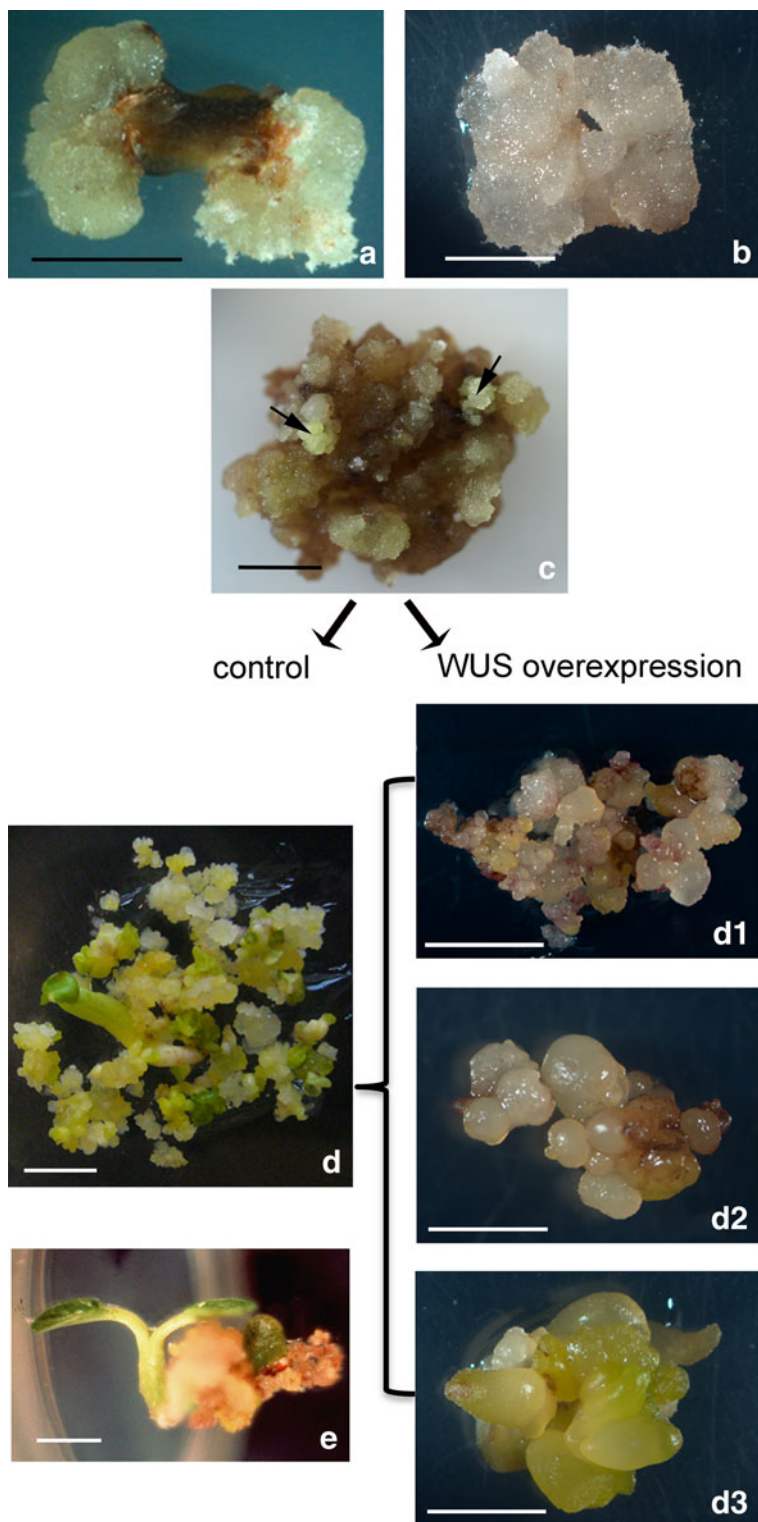
Cultures of *A. tumefaciens* were initiated from a single plated colony or from glycerol stocks and grown overnight at 28°C with shaking (150 rpm) in liquid Luria–Bertani medium containing 50 mg/L kanamycin, carbenicilin

and rifampicin, to mid-log phase ($\text{OD}_{660} = 0.9\text{--}1.2$). The *A. tumefaciens* cells were collected by centrifugation and resuspended in liquid inoculation medium same as the BM plant medium. The *A. tumefaciens* culture was then diluted 1/50 for explant inoculation. Hypocotyl fragments were dipped in the *A. tumefaciens* suspension culture for 25 min. The bacterial suspension was then removed and the explants were wiped on filter paper to remove excess bacteria. Explants were placed on callus induction medium (BM medium supplemented containing 2,4-D and kinetin at the concentration of 0.1 mg/L) at 25°C for 48 h for co-cultivation and then subcultured on the same callus induction medium supplemented with 25 mg/L of kanamycin and 500 mg/L of cefotaxime. Subcultures were performed every 14 days. Cefotaxime concentration was decreased gradually to 250 mg/L after the second subculture, 125 mg/L after the fourth subculture. The cultures were kept at $29^{\circ} \pm 1^{\circ} \text{C}$ under 16–8 h photoperiod.

Plasmid constructs

Four constructs were made using the pGWB Gateway cloning series (Nakagawa et al. 2007): pGWB2-*WUS* (35S:*WUS*), pGWB5-*WUS* (35S:*WUS*-GFP fusion), pGWB2-GFP (35S:GFP), and pGWB1 (empty vector: no promoter, no gene). pGWB2-GFP and pGWB1 were used as controls in transformation experiments. The pENTR-*WUS* cDNA (provided by Pr. Laux' laboratory, University of Freiburg, Germany) was cloned in the REGIA project (Paz-Ares and The REGIA Consortium 2002). The *WUS* cDNA was amplified from *A. thaliana* using specific primers *AtWUS*-1 (5'ATGGAGCCGCCACAGC) and *AtWUS*-2 (5'CATGTTCAGACGTAGCTC) and cloned into the pCR II TOPO blunt vector (Invitrogen). Subsequently, the *WUS* open reading frame (ORF) was amplified with the primers attB1*AtWUS* (5'AAGGGGACAAGTTTGTACAAAAAAGCAGGCTCCATGAGAGGATCCATGGTGAGCAA) and attB2*AtWUS* (5'AAGGGGACCCTTTGTACAAGAAAGCTGGGTTCATGTTCAGACGTAGCT) which were used to amplify *AtWUS* and recombined the gene into pDONR201 (Invitrogen). The pENTR-*GFP* was obtained by PCR amplification of *GFP* gene from the pGWB5 vector using attB1*GFP* (5'GGGGACAAGTTTGTACAAAAAAGCAGGCTCCATGAGAGGATCCATGGTGAGCAA) and attB2*GFP* (5'GGGGACCCTTTGTACAAGAAAGCTGGGTTCATGTTCAGACGTAGCT) and recombination into pDONR207 (Invitrogen). The pENTR-delta used for deleting gateway cassette from pGWB1 was obtained by modification of pENTR11 (Invitrogen). The *ccdB* gene for negative selection was deleted from pENTR11 by *SalI*–*XhoI* restriction enzyme digestion and religation. The inserts in pENTR-*WUS*, pENTR-GFP and pENTR-delta were transferred by LR recombination in the destination binary vectors

Fig. 1 Somatic embryogenesis in control and WUS-overexpressing tissues: **a** transformed calli on hypocotyl explant, **b** isolated callus, **c** embryogenic tissues (*arrows*) on callus. **d** Embryogenic lines: **d** control; **d1**, **d2**, **d3** successive stages of a WUS-overexpressing line, **e** somatic embryo development on control line (*bar* 0.5 cm)



pGWB2, pGWB5 and pGWB1, respectively. The fragment generated by PCR, the ligated junctions and the cloned fragments were verified by sequencing in all vectors. These plasmids were transferred into the *A. tumefaciens* strain C58::pGV2260 (Deblaere et al. 1985) by electroporation.

RT-PCR analysis

Total RNA was isolated from 100 mg samples with the RNeasy mini kit (QIAGEN S.A., Courtabeuf, France) following the manufacturer's instructions, and including the

optional RNase-free DNase step to avoid contamination with genomic DNA. RNA was extracted from embryogenic tissues from transformed and untransformed lines. For the transformed lines, successive stages of differentiation were studied in the transformed lines (see “Results”). Reverse transcription of mRNA was carried out in a 20 μ L final volume from 1 μ g total RNA with the SuperScript II reverse transcriptase (Invitrogen) according to manufacturer’s instructions. PCRs contained 2 μ L of cDNA, corresponding to 50 ng of total RNA, in a 50 μ L final volume, 1X PCR buffer, 0.2 mM dNTP, 1.5 mM MgCl₂, *Taq* DNA polymerase native and recombinant (Invitrogen®) 0.05 U and 0.2 μ M of each primer (*AtWUS-1* and *AtWUS-2* see above).

Histological analysis

Embryogenesis tissues were sampled at successive stages of differentiation and fixed in 0.2 M phosphate buffer at pH 7.2, supplemented with 2 % (v/v) paraformaldehyde, 1 % (w/v) caffeine, and 1 % (v/v) glutaraldehyde in a vacuum chamber for 30 min, then overnight at 4 °C. Tissues were dehydrated in ethanol, from 70 to 100 % progressively, then impregnated with ethanol–resin (50/50) for 2 h, and finally with resin 100 % overnight at 4 °C. Samples were embedded in Technovit 7100 resin (Heraeus Kulzer GmbH, Germany). Finally, 4- μ m sections were double-stained with periodic acid Schiff (PAS) (Merck) and Naphthol Blue Black (NBB) (Sigma-Aldrich) (Buffard-Morel et al. 1992), and imaged with a Leica DMRXB microscope.

Confocal and scanning electronic microscopy

The WUS–GFP signal was characterized by confocal laser scanning microscopy (Confocal LSM710 ZEISS CARL SAS) in whole mounts or fresh hand-cut 1-mm sections. Spectral analysis confirmed that the observed fluorescence corresponded to GFP and not autofluorescence. Vibratome sections of the samples expressing *AtWUS-GFP* were stained with 4,6-diamidino-2-phenylindole dihydrochloride (DAPI) at 1 g/L in PBS for 15 min, then rinsed and mounted in water. For scanning electronic microscopy, fresh tissues were cooled at –33 °C by a Deben Coolstage and observed with a Hirox SH-1500 benchtop SEM.

Determination of auxin content

Auxin (IAA) content was measured in cotton callus. The procedure used homogenized frozen tissue. Samples were weighted and 10 ng of a standard ¹³C₆-IAA (Cambridge Isotope Laboratory Inc.) was initially added as internal tracers for recovery and analytical purposes used to quantify AIA as described previously (Denancé et al. 2012).

Statistical analyses

Induction of somatic embryogenesis was quantified through a logistic regression with the R software environment for statistical computing and graphics (www.r-project.org/).

Comparison of IAA content in calli expressing WUS and in control calli was performed with a non-parametric test (Kruskal and Wallis) using Rcmdr package of R software (<http://cran.r-project.org/web/packages/Rcmdr/>).

Results

We have analyzed the effect of *WUSCHEL* overexpression during regeneration via somatic embryogenesis, from the appearance of embryogenic tissues through the development of somatic embryos. *AtWUS*-expressing tissues were compared to controls with regard to developmental and morphological characteristics. Figure 1 summarizes the successive steps of the regeneration process for both control and *AtWUS*-expressing calli.

WUSCHEL enhances the induction of somatic embryogenesis

In six independent experiments, each including ~100 explants per construct, we observed a significant increase in the percentage of embryogenic lines in calli where *WUS* was overexpressed (Table 1). *AtWUS* overexpressors yielded 3–4 times more embryogenic lines than control *GFP* overexpressors. Variability between experiments probably reflects the heterogeneity of the starting material. For example, experiments 1 and 2 were done with a different seed batch than experiment 3.

The positive effect of *AtWUS* overexpression was confirmed in another experiment where pGWB1 was used as the second control (repetition 7 in Table 1). This result shows that, the *GFP* gene has no detrimental effect on somatic embryogenesis. The percentage of explants giving rise to embryogenic tissues was statistically higher when the fusion *AtWUS-GFP* is overexpressed compared to *AtWUS*. In some cases, we have observed the appearance of embryogenic calli directly on the explant expressing the fusion *AtWUS-GFP* (Fig. 2). Compared to the *GFP* control, *AtWUS* overexpressor calli produced many more embryogenic cell clusters (Fig. 3a, b). The clusters contained dark blue-stained active cells, indicating a high rate of soluble proteins, with a thick cell wall, a big nucleus and a single nucleolus (Fig. 3b'). These cytological features are common to embryogenic cells in general whatever the species (Michaux-Ferrière and Schwendiman 1992).

Table 1 Induction of somatic embryogenesis

	pGWB1			pGWB2GFP			pGWB2WUS			pGWB5WUS		
	Explants	Emb calli ^a	% Explant giving Emb calli	Explants	Emb calli ^a	% Explant giving Emb calli	Explants	Emb calli ^a	% Explant giving Emb calli	Explants	Emb calli ^a	% Explant giving Emb calli
Repetition 1				81	5	6.17	79	14	17.7			
Repetition 2				72	10	13.9	102	15	14.7			
Repetition 3				120	32	26.7	117	62	53			
Repetition 4				113	13	11.5				123	24	19.5
Repetition 5				114	1	0.9				120	16	13.3
Repetition 6				56	10	21.3				98	74	77.1
Repetition 7	112	12	10.7	112	8	7.2	150	42	28	128	66	51.6
Total	112	12	10.7	668	79	11.8	448	133	29.7	469	180	38.4

^a Number of explants giving rise to embryogenic calli. If an explant gave several embryogenic lines, it was counted as one

Embryogenic lines overexpressing *AtWUS* show specific morphological and histological features

The macroscopic morphology of embryogenic lines overexpressing *GFP* was identical to that of lines we previously transformed with *GUS* gene or various genes of interest (more than five different types). The tissues overexpressing *GFP* have passed through the classical stages leading to somatic embryogenesis and plantlet regeneration, typically observed in cotton (Fig. 1). Different stages of differentiation are usually found in a single cotton embryogenic line and the embryogenic lines can simultaneously proliferate and form embryos during several years on hormone-free medium (Fig. 1d). Accordingly, control embryogenic lines, including those overexpressing *GFP*, produced several types of structures: aggregates of embryogenic cells, pro-embryos and embryos at different stages of development (Fig. 1d, e). Embryogenic lines overexpressing *AtWUS* differed from this typical scenario and produced tissues

going through three distinct stages of differentiation. The first stage (S1) corresponded to embryogenic aggregates same as control at the time of appearance but that rapidly evolved in clusters, initially forming small pale yellow round structures (Fig. 1d1), then growing into large round structures distinguished as the second stage (S2) (Fig. 1d2). In the third stage (S3), these structures evolved into even larger green masses, usually not observed during cotton regeneration (Fig. 1d3). At stage 3 and beyond, these structures can exhibit characteristics of somatic embryos with a well-organized root pole (Fig. 4c). We conclude that the formation of abnormal embryo-like structures results from the constitutive expression of *AtWUS*.

Tissues in embryogenic lines overexpressing either *AtWUS* or *GFP* were conducted. Similar to control embryogenic tissues, the *AtWUS*-induced S3 embryo-like structures can be maintained on a hormone-free medium, through multiplication of similar adventitious S3 embryos produced by “budding” (Fig. 1d3). More than 100 embryogenic lines were thereby subcultured with a stable phenotype for over 2 years.

AtWUS ectopic expression promotes the formation of leaf-like structures

Despite the fact that embryoids over expressing *WUSCHEL* have never gone into differentiation of shoots, ectopic leaf-like structures were produced by these embryoid formations (Figs. 3c, 4). This phenomenon has never been observed on control somatic embryos. Histological examination showed that these ectopic leafy structures arise from the peripheral zone of the embryo-like structures where cells actively divide (Figs. 3c, d, 5e, f). The leaf-like formations developed on embryos expressing *AtWUS* alone as well as *AtWUS-GFP*. In the latter case, the corresponding *GFP* fluorescence was

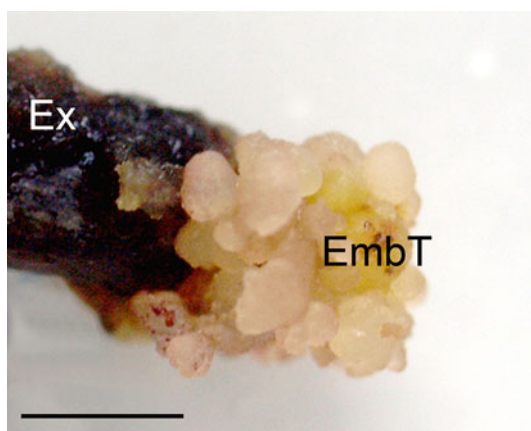


Fig. 2 Direct embryogenesis on hypocotyl explant expressing *WUS-GFP*. *Ex* explant, *EmbT* embryogenic tissue (bar 0.5 cm)

Fig. 3 Histological examination (NBB staining) of *WUS*-induced embryogenic line (EmbS) and embryo-like structures (ES). **a** Control *GFP*-expressing EmbS, **b** *WUS*-expressing EmbS with multiple embryogenic cell clusters (arrows), **b'** detail of embryogenic cells, **c** *WUS*-expressing ES showing ectopic development of bud-like formation, **d** detail showing that ectopic formations originated from the peripheral zone. Bar **a, b** 150 μ m; **c, d** 500 μ m

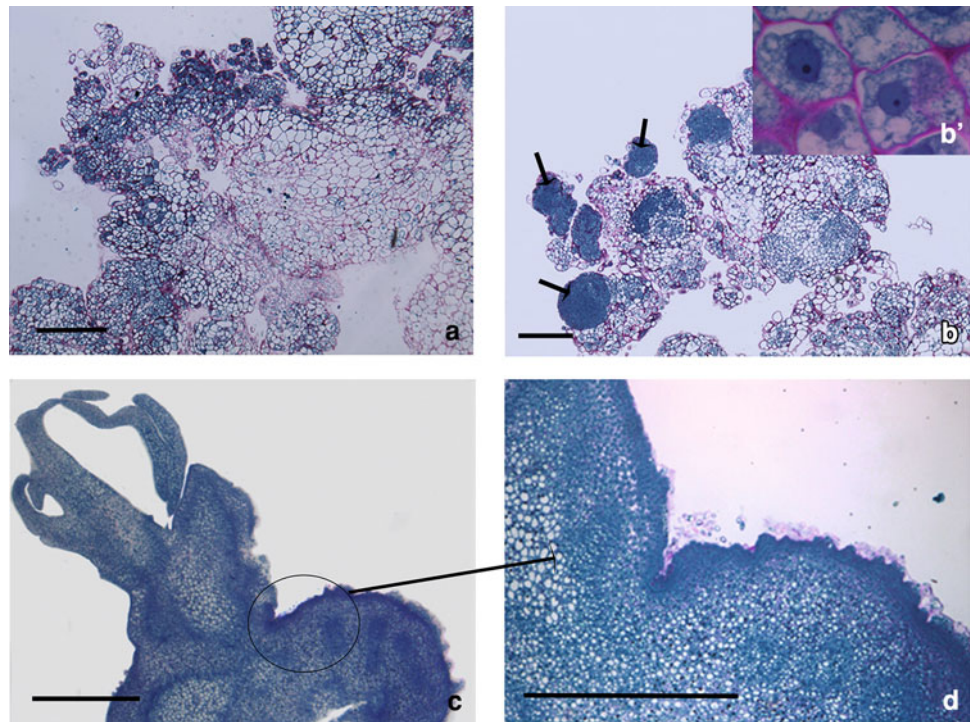
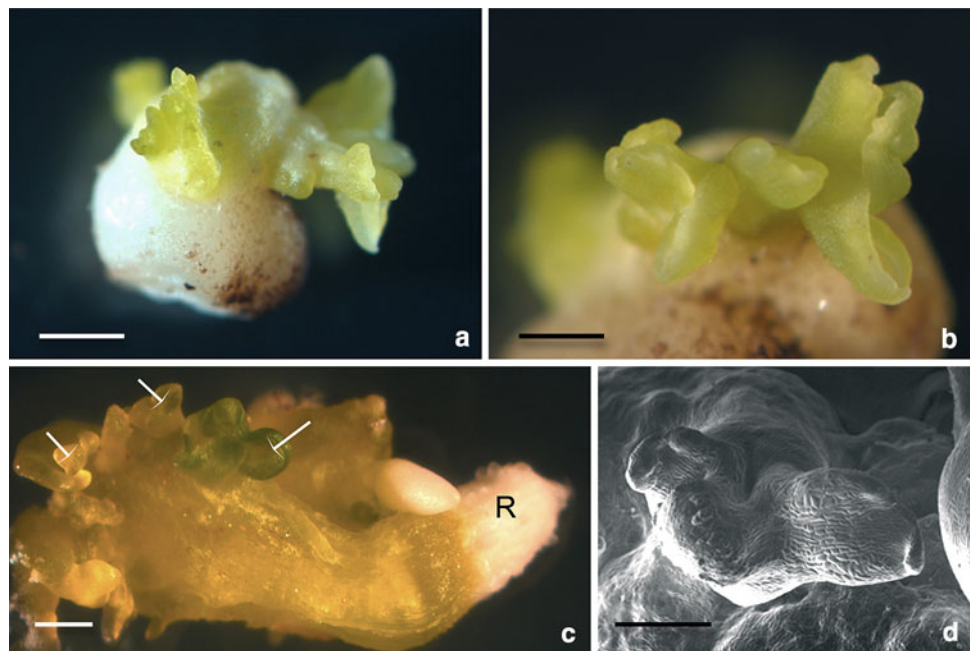


Fig. 4 **a–c** Embryo-like structures supporting ectopic organogenesis from embryogenic lines overexpressing *WUS*; **d** MEB picture of leaf-like formations. *R* root; arrows point to leaf-like formations (bar 1 mm)

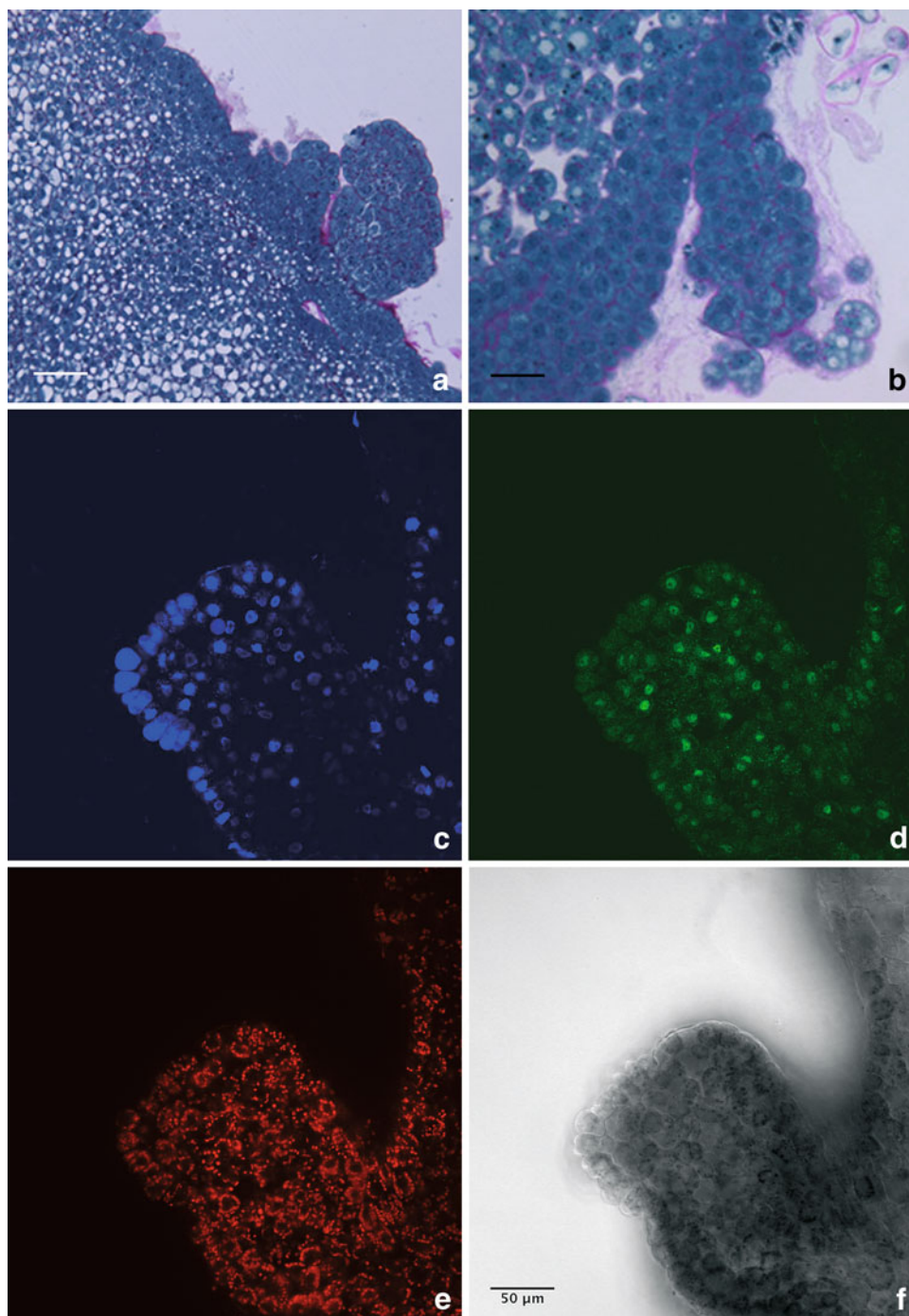


detected in the nucleus as expected for the *AtWUS* fusion protein. Confocal imaging of S3 embryogenic-like formations expressing *AtWUS-GFP* revealed that signal was highest in the globular formations arising at the peripheral zone (Fig. 5c, d). These formations are constituted of active meristematic cells as shown by NBB coloration (Fig. 5a, b). This observation suggests that the expression of *AtWUS* leads to the reactivation of cells giving rise to organogenic structures.

Embryo-like formations depend on *AtWUS* expression

Some of the embryogenic lines transformed with the *AtWUS* or *GFP-AtWUS* transgenes did not show a developmental pattern giving rise to embryo-like structures. Instead, they only grew as proliferating tissues without any embryonic differentiation, remaining at stage S1. Therefore, we tested the *WUSCHEL* expression in this type of tissue as well as in the S3 structures. RT-PCRs have been

Fig. 5 **a, b** Budding structures emerging from the peripheral zone of the embryo-like structures and cellular content of these type of formations arising from *WUS*-induced embryo-like structures (NBB stained). **c** DAPI staining, **d** *WUS*-GFP fluorescence, **e** chlorophyll fluorescence (*bar* 50 μm)



done with tissues exhibiting globular green masses (Fig. 1d3) and cultures of undifferentiated *WUS* lines. All tissues were sampled after the same delay of 1 year in *in vitro* culture. *AtWUS* expression was only detected in embryogenic S3 structures (Fig. 6). The results indicate that transformed embryogenic tissue not evolving into S2 and S3 embryo-like structures do not express *AtWUS*, confirming that S3 formations and ectopic leaf-like structures result from overexpression of *AtWUS*.

WUSCHEL did not interact with endogenous IAA content

Endogenous IAA levels were measured in *AtWUS* overexpressing embryogenic callus and in *GFP* overexpressors, for two independent experiments with ten samples. No differences were seen for IAA content between *AtWUS* ($1.68 \text{ pg} \cdot \text{mg}^{-1}$) and *GFP* ($1.31 \text{ pg} \cdot \text{mg}^{-1}$) expressing lines (Kruskal–Wallis rank sum test, p value = 0.32).

Discussion

With this work, we wished to test whether genes that stimulate organogenesis or somatic embryogenesis in *A. thaliana* promote the same developmental programs in cotton tissues and can induce the regeneration process starting with explants from recalcitrant cultivars.

In our hands and with our standard in vitro culture protocol, the overexpression of *AtWUS* in tissues of a recalcitrant variety (a CIRAD-IRAD variety, Irma96 + 97) did not result in the production of regenerated plants. This achievement will still require the optimization of other non-genetic factors, such as the balance of exogenous phytohormones and the composition of in vitro culture media. Nevertheless, we showed that *AtWUS* overexpression in an in vitro routinely used genotype improved somatic embryogenesis and induced organogenesis on embryo-like structures cultured on a hormone-free medium.

The positive effect of *AtWUS* on somatic embryogenesis was observed in all experiments, when overexpressed by itself or as a translational fusion with *GFP*. However, the fraction of explants giving rise to embryogenic tissues was higher with *AtWUS-GFP*, possibly because the fusion product may be more stable.

Embryogenic lines overexpressing *AtWUS* evolved following a quite different scenario from the one we usually observe in cotton. In the classical scenario, different stages

of differentiation are usually found in a single embryogenic line: as soon as embryogenic tissues are observed on calli they can be isolated on a hormone-free medium. On this medium, the embryogenic culture can proliferate for years by regular subcultures. Two phenomena are observed simultaneously: proliferation of embryogenic clumps and somatic embryo differentiation.

In a *AtWUS* overexpressing context and without exogenous phytohormones, three distinct stages were observed leading to large embryo-like green masses, highly differentiated and supporting adventitious organogenesis. Histological examinations showed that, in contrast to control lines, *AtWUS* overexpressing lines start to differentiate very actively. Highly active zones, defined as clusters of embryogenic cells, appeared only in the *WUS* tissues. In the third stage of development, the highly active cells were located in the peripheral zone of the embryo-like structures. The active zones overlap with *AtWUS* expression as confocal examination of lines expressing *AtWUS-GFP* fusion showed fluorescence in the corresponding zones. These peripheral active zones give rise to adventive bud-like formations. *AtWUS* overexpression triggered the cell totipotency in these tissues and lead to new meristems. Xu et al. (2005) observed similar results where in their system, ectopic flower meristems were initiated from the differentiated cortex cells. The reason why *AtWUS* expression is limited or at least stronger to the peripheral zone, while

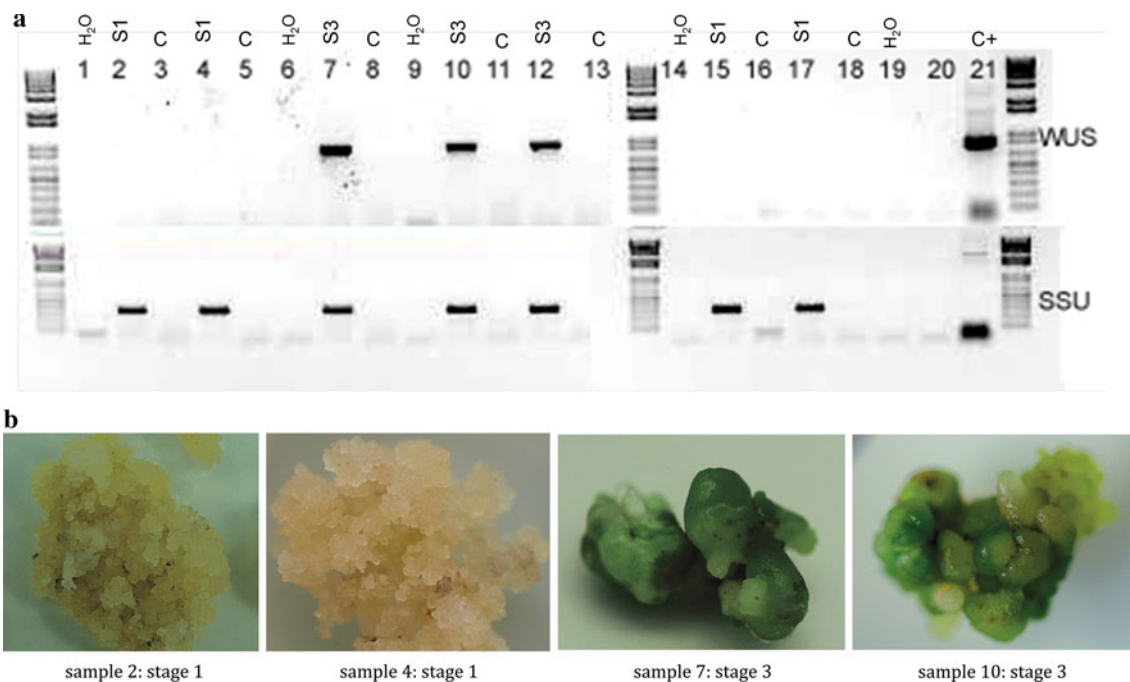


Fig. 6 Analysis of *WUS* ectopic expression in transgenic tissues. **a** RT-PCR analysis of *WUS* transcript level. *S1* cultures at stage 1, *S3* cultures at stage 3; *C* RNA control to show that the amplification is not due to residual DNA; *C+* positive *WUS* control. *Bottom fragment*

corresponds to amplification of the small ribosomal subunit (SSU) cDNA. **b** Macroscopic views of the cultures, at stages 1 and 3, analyzed by RT-PCR

under the 35S promoter it is supposed to be overexpressed in all tissues, is unknown. *AtWUS* expression is probably defined by the distribution of *CLV3* signaling peptides as *CLAVATA* and *WUS* are known to be regulated in a feedback loop (Schoof et al. 2000; Brand et al. 2000).

Considering the new knowledge in hormonal control of shoot stem cell niche in interaction of transcription factors (Leibfried et al. 2005; Zhao et al. 2010), we can say that cell differentiation observed in *WUS* over-expressors is due to an interaction of *WUS* with phytohormones. In our conditions, *WUSCHEL* did not alter IAA activity as no differences were seen for IAA content between *WUS* and *GFP*-expressing lines whilst on the other hand *WUS* expression facilitates high cytokinin activity in the SAM (Leibfried et al. 2005; Shani et al. 2006).

We demonstrate in this work that *AtWUS* overexpression dramatically promotes the production of embryogenic lines and could potentially improved regeneration. We tried to regenerate plants from these tissues obtained in every transformation experiments performed and could potentially improve plant regeneration. We have regenerated few plants from *AtWUS* overexpressing tissues from our experiments. These plants had present a strong *WUS* phenotype (Kieffer et al. 2006; Xu et al. 2005) with many branches and waffle-curved leaves and expressed the *Arabidopsis* gene at detectable levels from tissues taken at several parts of the plants (data not shown). Therefore, we speculate that the *Arabidopsis WUS* gene is useful to improve the regeneration/transformation process in cotton but only if it is induced at critical steps, for example, when in vitro regeneration is initiated. Separate studies focusing on the effect of *AtWUS* ectopic expression in pepper (Solis-Ramos et al. 2009) and coffee (Arroyo-Herrera et al. 2008) under the control of estradiol-inducible promoter resulted in no normal plant regenerated because of the leakiness of the promoter. In White spruce (Klimaszewska et al. 2010), very few and severely abnormal *WUS* transgenic somatic embryos developed on medium containing 17- β -estradiol and morphologically normal somatic embryos were collected only on media containing no, or very few concentrations of the inducer. In future studies, other types of inducible systems should be tested to find a reliable one that insure expression of *WUS* transgene only at specific steps of regeneration process.

Our results have shown that the overexpression of the *WUS* gene from *Arabidopsis* can promote the expression of embryogenic competence of dedifferentiated proliferating cells obtained on cotton hypocotyl explants. Further experiments are needed to understand interactions between endogenous hormone and *WUS* expression to use *WUS* over-expression to regenerate recalcitrant genotypes.

We can notice that our approach could lead to a promising way to obtain marker-free transgenic plants. Using an

Agrobacterium binary vector carrying *WUS* and a gene of interest, the transformed calli would be the only one able to give rise to embryogenic lines and plants. A co-transformation method or the use of two binary vectors in a same *Agrobacterium* would allow avoiding the presence of “embryogenic gene” in plants through subsequent segregation.

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