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TECHNICAL ADVANCE

Isolation of precise plastid deletion mutants by homology-based excision: a resource for site-directed mutagenesis, multi-gene changes and high-throughput plastid transformation

Vasumathi Kode, Elisabeth A. Mudd, Siriluck lamtham[†] and Anil Day^{*}

Faculty of Life Sciences, The University of Manchester, 3.614 Stopford Building, Oxford Road, Manchester M13 9PT, UK

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^{*}For correspondence (fax +44 161 275 3938; e-mail anil.day@manchester.ac.uk).

[†]Present address: Kasetsart University, Kamphaengsaen Campus, Nakorn Prathom 73140, Thailand.

Summary

We describe a simple and efficient homology-based excision method to delete plastid genes. The procedure allows one or more adjacent plastid genes to be deleted without the retention of a marker gene. We used *aadA*-based transformation to duplicate a 649 bp region of plastid DNA corresponding to the *atpB* promoter region. Efficient recombination between *atpB* repeats deletes the intervening foreign genes and 1984 bp of plastid DNA (co-ordinates 57 424–59 317) containing the *rbcl* gene. Only five foreign bases are present in $\Delta rbcl$ plants illustrating the precision of homology-based excision. Sequence analysis of non-functional *rbcl*-related sequences in $\Delta rbcl$ plants indicated an extra-plastidic origin. Mutant $\Delta rbcl$ plants were heterotrophic, pale-green and contained round plastids with reduced amounts of thylakoids. Restoration of autotrophy and leaf pigmentation following *aadA*-based transformation with the wild-type *rbcl* gene ruled out mutations in other genes. Excision and re-use of *aadA* shows that, despite the multiplicity of plastid genomes, homology-based excision ensures complete removal of functional *aadA* genes. Rescue of the $\Delta rbcl$ mutation and autotrophic growth stabilizes transgenic plastids in heteroplasmic transformants following antibiotic withdrawal, enhancing the overall efficiency of plastid transformation. Unlike the available set of homoplasmic knockout mutants in 25 plastid genes, the *rbcl* deletion mutant isolated here is readily transformed with the efficient *aadA* marker gene. This improvement in deletion design facilitates advanced studies that require the isolation of double mutants in distant plastid genes and the replacement of the deleted locus with site-directed mutant alleles and is not easily achieved using other methods.

Keywords: Nupts, photosynthesis, ribulose biphosphate carboxylase/oxygenase, targeted gene deletion, tobacco.

Introduction

The simplicity and precision of homology-based excision, together with the predominance of homologous recombination in plastids, makes it a particularly attractive tool for manipulating plastid genomes. Use of native plastid enzymes avoids the extra transformation steps needed to introduce foreign site-specific recombinases, unintended site-specific recombinase-promoted rearrangements and

the retention of site-specific recombinase target sites in engineered plastid genomes (Corneille *et al.*, 2001, 2003; Hajdukiewicz *et al.*, 2001; Lutz *et al.*, 2004, 2006). Homology-based excision has allowed the removal of marker genes from transgenic plastids (Fischer *et al.*, 1996; lamtham and Day, 2000; Klaus *et al.*, 2004). However, the potential of exploiting native plastid recombination pathways to

manipulate plastid DNA has not been fully explored. To improve the design of deletion mutants to facilitate advanced studies on plastid genes, we examine whether homology-based excision can be used to make precise deletions in plastid DNA without a requirement for a marker gene to select mutant plastids.

Loss-of-function plastid mutants resulting from precise targeted deletion of plastid genes without the insertion of marker genes have not been described in flowering plants. All described knockout mutations in 29 plastid genes contain targeted insertions of the *aadA* marker gene used to select transformed plastids (reviewed by Maliga, 2005). Marker-free plastid mutants would provide new opportunities and advantages compared to the use of conventional *aadA* insertion mutants and wild-type (WT) plants in transplastomic research. They can be transformed with *aadA*, which is the most widely used plastid transformation marker gene (Svab and Maliga, 1993) due to the relative inefficiency of alternative markers (Carrer *et al.*, 1993; Huang *et al.*, 2002; Svab *et al.*, 1990). When a modified plastid gene containing a site-directed mutation is transformed into a deletion mutant, all transformants will contain the modified allele whose phenotype will not be masked by the resident wild-type allele. The introduction of site-directed mutations into WT plastids can be problematic due to undesirable recombination events in the region between marker and mutation (Andrews and Whitney, 2003; Whitney *et al.*, 1999). Marker-free mutants facilitate the stepwise introduction of mutations into the plastid genome to isolate plants with mutations in several plastid genes. Co-transformation with two plasmids provides a less predictable method for integrating changes at multiple sites within the plastid genome (Carrer and Maliga, 1995; Ye *et al.*, 2003). Efficient *aadA*-based antibiotic selection combined with mutant rescue using the wild-type plastid allele (Klaus *et al.*, 2003) enables rapid identification of plastid transformants and stabilizes recombinant genomes in transformed plants. This provides the basis for high-throughput *aadA*-based transformation of foreign trait genes into plastids. Pleiotropic effects of marker gene expression on transcription or protein accumulation within mutant plastids are removed, and marker-free plastid mutants are likely to satisfy regulatory approval for widespread dissemination and use of plastid mutants within the research community.

Here we target the *rbcl* gene, which encodes the large subunit (LS) of ribulose biphosphate carboxylase/oxygenase (RuBisCO), the key enzyme in photosynthetic carbon assimilation (Andrews and Whitney, 2003; Parry *et al.*, 2003). Our procedure is simple, can be used to delete one or more adjacent plastid genes, and is applicable to non-essential plastid genes, whose mutation allows growth on media containing sucrose.

Results

Plastid transformation vectors and experimental design

The approach we used to delete the *rbcl* gene is illustrated in Figure 1(a). In vector pUM83, the *aadA* marker gene is placed downstream of the *rbcl* gene. Sequences upstream of *rbcl* are duplicated and placed downstream of *rbcl* and *aadA*. This creates a direct repeat flanking *rbcl* and *aadA*. Recombination between these direct repeats excises the *rbcl* and *aadA* genes in one event. The advantage of this strategy is that all mutant $\Delta rbcl$ cells will also be free of the *aadA* marker gene. This means that a visual screen for plastid mutants, which often have a pigment-deficient phenotype (Kanevski and Maliga, 1994; Klaus *et al.*, 2003; Swiatek *et al.*, 2003), will also identify *aadA*-free plants.

Vector pUM83 (Figure 1a) contains converging expression cassettes with the *aadA* marker (Goldschmidt-Clermont, 1991) and *uidA* reporter genes (Jefferson, 1987) inserted into the *Aocl* site located in the intergenic region between the *rbcl* and *accD* genes (Wakasugi *et al.*, 2001). The 5' *atpB* regulatory region drives expression of *uidA* and is comprised of 649 bases including the ATG initiation codon. This creates two 649 bp direct repeats, DR1 and DR2 (Figure 1a), comprised of 5' *atpB* regulatory sequences that flank the *rbcl* gene. The 5.7 kbp left and 2.1 kbp right plastid (pt) DNA sequences bordering *aadA* and *uidA* target integration by homologous recombination following plastid transformation. Transgenic plastid genomes are then selected with antibiotics to replace WT plastids. A recombination event between the 649 bp direct repeats will excise the *rbcl*, *aadA* and *uidA* genes as a 6.1 kbp circle with a single *HindIII* site, leaving a 153 kbp marker-free $\Delta rbcl$ plastid genome (Figure 1a). The 6.1 kbp circle lacks sequences necessary for stable maintenance as an episome and is lost. In the absence of selection, marker-free $\Delta rbcl$ plastid genomes will accumulate, leading to the isolation of marker-free mutant plants.

Isolation of pUM83 (T_0) plastid transformants selected with antibiotics

Plastid transformation was carried out using particle bombardment on leaf explants (Kode *et al.*, 2005). Green resistant shoots were taken through three rounds of regeneration on media containing spectinomycin and streptomycin. Of seven antibiotic-resistant clones isolated, three stained positive for GUS using X-gluc (not shown), indicating the presence of an active *uidA* gene. Two of these GUS-positive clones (8a and 8c) were taken forward for further analysis. DNA blot analysis indicated replacement of WT ptDNA with transgenic plastid genomes (see below). Seeds were collected from selfed T_0 pUM83 transplastomic plants.

Figure 1. Plastid DNA recombination events underlying deletion and restoration of *rbcL*.

(a) Scheme showing isolation of $\Delta rbcL$ plastid (pt) DNA. Left (map co-ordinates 53 608–59 318) and right (59 319–60 859) ptDNA sequences in vector pUM83 target integration of *aadA* and *uidA* into WT ptDNA by homologous recombination. Recombination between 649 bp direct repeats DR1 and DR2 comprised of the *atpB* ATG initiation codon and upstream region excises a 6.1 kbp circle with foreign and *rbcL* genes to leave a marker-free $\Delta rbcL$ plastid genome. DR1, DR2 and the DR1+2 products of recombination are identical.

(b) Integration pattern of the pUM79 vector. Shown are *Hind*III (H) sites and fragments that hybridize to *rbcL*, *aadA* and *accD* probes. Plastid *rbcL*, *accD*, *psaI* and *ycf4* genes and foreign genes are indicated. Hybridization probes and *aadA* F + R primers are located. The WT ptDNA *Hind*III site at 63 894 bp is indicated as '63 864' in recombinant and $\Delta rbcL$ ptDNAs.

Isolation of $\Delta rbcL$ mutant plants (T_1 generation)

Seeds from pUM83-8a and -8c transplastomic parents were germinated on MS medium containing spectinomycin (500 mg l^{-1}) or MS medium without antibiotics. All of the 30 pUM83-8a (T_1) seedlings were green and resistant to spectinomycin (not shown). Two white sensitive seedlings were found amongst the 30 T_1 progeny of pUM83-8c grown on spectinomycin-containing medium (Figure 2a) and resembled spectinomycin-bleached WT seedlings. On antibiotic-free medium, three pale-green pUM83-8c seedlings

(Figure 2b) were found amongst 30 T_1 seedlings and were propagated *in vitro* for further analysis. On further growth on MS medium without antibiotics, half of the T_1 seedlings from pUM83-8a and -8c plastid transformants gave rise to plants with one or more true leaves containing pale-green sectors (Figure 2c). WT plants do not contain pale-green sectors (Figure 2d). T_2 seeds were collected from variegated T_1 plants, allowing long-term storage of the mutant (Figure 2e,f).

A pale-green phenotype is characteristic of mutations in some photosynthesis-related genes (Klaus *et al.*, 2003;

Swiatek *et al.*, 2003) including *rbcl* (Kanevski and Maliga, 1994). Sensitivity to spectinomycin and a pale-green phenotype would result from excision of the *aadA* and *rbcl* genes (Figure 1). To purify pale-green mutant plants, three rounds of regeneration were performed on media lacking antibiotics using either leaves from pale-green plants (clone 8c) or pale-green sectors from variegated plants (clone 8a). Pale-green mutant plants (Figure 2g) were propagated by sub-culturing shoots on MS medium supplemented with 3% w/v sucrose.

PCR analysis of $\Delta rbcl$ mutant plants and sequencing confirms the precision of homologous recombination

Recombination within the duplicated *atpB* promoter region excises *rbcl* and the foreign genes. The precision of the recombination event was analysed by PCR using primers pt F and pt R (Figure 3c). The primers amplified a 3.2 kbp band from WT DNA (Figure 3a) and a 1.3 kbp band from pale-green $\Delta rbcl$ -8a plant DNA (Figure 3a). The absence of a WT 3.2 kbp band in $\Delta rbcl$ lanes is consistent with the absence of WT ptDNA but might also reflect preferential amplification of the smaller 1.3 kbp band. The 1.3 kbp amplification product from $\Delta rbcl$ plants was purified and sequenced. The sequence (Figure 3c) confirmed that the 1.3 kbp PCR product was the result of recombination between the 649 bp direct repeats (Figure 1a). No changes were found in the recombined copy of the 649 bp *atpB* 5' region remaining in $\Delta rbcl$ plants, demonstrating a perfect homologous recombination event between the two repeats. An introduced *NotI* site spans the junction between the deletion end points located at bases 57 423 and 59 318. The $\Delta rbcl$ plastid genome contains only 5 bp of foreign sequence (GGCCG) corresponding to internal bases in the *NotI* site (Figure 3c). The $\Delta rbcl$ plants should lack the *aadA* marker gene. Primers directed against *aadA* (see locations in Figure 1) amplified a band of the expected size from a uniformly dark-green pUM83-8a (T_0) plant DNA but not from WT or pale-green $\Delta rbcl$ -8a extracts (Figure 3b).

DNA blot analyses of $\Delta rbcl$ mutant plants

Total DNA was extracted from WT plants, a uniformly dark-green pUM83-8a (T_0) plant and two $\Delta rbcl$ mutant clones (8a,

Figure 2. Isolation and phenotype of $\Delta rbcl$ plants.

(a) pUM83 transplastomic 8c (T_1) seedlings on MS medium with spectinomycin (500 mg l^{-1}), white seedlings arrowed; (b) pUM83 transplastomic 8c (T_1) seedlings on MS medium, pale-green seedlings arrowed; (c) pUM83 transplastomic 8a (T_1) plant with sectors on MS medium; (d) WT plant growing on MS medium; (e, f) pUM83 8a (T_1) transplastomic plants growing in soil; (g) $\Delta rbcl$ plants growing on MS medium; (h) plastids (pt) in $\Delta rbcl$ leaves, magnification 11 000x, plastids (pt), stroma lamellae (L), grana (Gr); (i) WT and $\Delta rbcl$ shoots placed on RMOP media lacking sucrose for 3 weeks; (j) $\Delta rbcl$ leaves in X-gluc buffer showing GUS-negative phenotype; (k) antibiotic-resistant plant from $\Delta rbcl$ leaves transformed with pUM79; (l) leaves from pUM79 transformant in X-gluc buffer showing GUS-positive phenotype.

Figure 3. PCR analysis on pUM83 transplastomic and $\Delta rbcL$ DNA. Sources of plant DNA are indicated above lanes. Marker sizes are shown on the left of each gel photograph. (a) pt F and R primers flank the *rbcL* deletion and amplify a smaller product from $\Delta rbcL$ plant extracts; 1% w/v agarose gel; (b) *aadA*-F and R primers (located in Figure 1a) fail to amplify a product using WT and $\Delta rbcL$ -8A extracts, 2% w/v agarose gel; (c) location of primers on WT and $\Delta rbcL$ ptDNA. The binding site for pt F1 is not present in $\Delta rbcL$ ptDNA. Sizes of expected PCR products and the sequence at the junction of the *rbcL* deletion are shown.

8c) from independent pUM83 transformants 8a and 8c. The locations of *rbcL*, *accD* and *aadA* hybridization probes are shown in Figure 1(a). An *rbcL* probe hybridizes to an 11.5 kbp *Hind*III band in digests of WT DNA (Figure 4a, lane 1). This is replaced by two bands of 7.0 and 6.1 kbp in digests of leaf DNA from a T_0 pUM83-8a plant (Figure 4a, lane 2). The 7.0 kbp band corresponds to the recombinant plastid genome resulting from integration of *aadA* and *uidA* expression cassettes (Figure 1a). The less intense 6.1 kbp *Hind*III band corresponds in size to the predicted excision product containing foreign genes and *rbcL* that has a single

*Hind*III site (Figure 1a) and also hybridizes to *aadA* (Figure 4d, lane 2). A discrete low molecular weight band of uncut DNA from a T_0 pUM83-8a plant (Figure 4b, lane 2), located well below the shear size of the bulk of DNA (>20 kbp), might correspond to the 6.1 kbp excision product. No low molecular weight species were found in other lanes (Figure 4b).

No *rbcL* hybridization was expected to digests of $\Delta rbcL$ plant DNA, but we found a weakly hybridizing 11.5 *Hind*III band (Figure 4a, lanes 3 and 4) that co-migrates with the strong WT band (Figure 4, lane 1). This faint 11.5 kbp band is unlikely to represent residual copies of WT plastid genomes because WT ptDNA in mutant plants would be expected to give rise to green sectors. No dark-green sectors have been observed in $\Delta rbcL$ mutant plants propagated for three years on MS media. This indicates that the *rbcL* related sequences in $\Delta rbcL$ plants are not functional. Primers pt F1 and pt R

Figure 4. DNA blot analyses on pUM83 transplastomic (T_0), $\Delta rbcL$ mutant (T_1) and pUM79 transformants of $\Delta rbcL$ plants.

Sources of total DNA preparations are indicated above lanes. DNA was either digested with *Hind*III (a, c, d) or untreated (b) before loading on 0.8% w/v agarose gels. Blot wash conditions: $0.1 \times$ SSC at 60°C . Bands marked with an asterisk are derived from the 6.1 kbp episome. Hybridization probes (located in Figure 1a) are indicated below each panel. Linear dsDNA size standards (left) and hybridizing band sizes (right) are shown beside the panels.

Figure 5. Alignment of WT ptDNA sequence (co-ordinates 59 231–59 633) with $\Delta rbcl$ PCR product using primers pt F1 and pt R (see Figure 3c).

(locations shown in Figure 3c) should amplify residual copies of any WT DNA present in $\Delta rbcl$ plants because primer pt F1 binds to the end of the *rbcl* gene, which is deleted in $\Delta rbcl$ plants. A 0.8 kbp PCR product amplified from $\Delta rbcl$ DNA contained seven base substitutions and six insertion/deletion events in a 386 bp region when compared to WT ptDNA (Figure 5). This sequence divergence is typical of plastid DNA sequences transposed to the nucleus (Huang *et al.*, 2005). No WT *rbcl* sequences were detected in $\Delta rbcl$ plants.

Hybridization of the *accD* probe to *Hind*III digests revealed an 11.5 kbp WT band (Figure 4c, lane 1), an 8.7 kbp band in pUM83 transformants (Figure 4c, lane 2), and a 9.6 kbp band in $\Delta rbcl$ plants (Figure 4c, lanes 3 and 4) as expected (see Figure 1a). An *aadA* probe hybridizes to expected bands of 8.7 and 6.1 kbp in pUM83 transformants (Figure 4d, lane 2) but did not hybridize to digests of WT and $\Delta rbcl$ DNA (Figure 4d, lanes 1, 3, 4); refer to maps in Figure 1(a) for sizes. Blot analyses confirm the isolation of *aadA*-free $\Delta rbcl$ mutant plants.

Phenotype of $\Delta rbcl$ plants

Sections of pale-green $\Delta rbcl$ leaves contained a predominance of circular plastids (pt) in which stroma lamellae (L) and grana (Gr) were clearly visible (Figure 2h). Much of the stroma was devoid of grana and stroma lamellae, unlike the typical lens-shaped chloroplasts found in leaf mesophyll cells in WT plants (see examples of WT plastids in Kode *et al.*, 2005; Swiatek *et al.*, 2003). Shoots formed on RMOP regeneration medium supplemented with 3% w/v sucrose were transferred to RMOP medium lacking sucrose. WT shoots continued to grow on RMOP medium lacking sucrose while $\Delta rbcl$ shoots from clone 8a died (Figure 2i). This indicates that mutant $\Delta rbcl$ plants are heterotrophic and

unable to grow on media lacking sucrose. The large subunit (LS) of RubisCO was not detected in $\Delta rbcl$ mutant plants by Western blot analysis (not shown).

Repair of the lesion in $\Delta rbcl$ mutant plants by transformation with the WTgene

Transformation of $\Delta rbcl$ plants with plastid transformation vector pUM79 introduces *aadA* and the WT *rbcl* allele into mutant plastids (Figure 1b). The *Hind*III restriction patterns of recombinant ptDNA in pUM79 and pUM83 transformants are different (see Figure 1a,b). This allows us to distinguish pUM79 transformants from the primary pUM83 transformants from which the $\Delta rbcl$ plants, used as recipients of transformation, were derived. Spectinomycin-resistant shoots with either dark-green or pale-green leaves were isolated 4–8 weeks after transformation. PCR analyses with *aadA* primers indicated the presence of *aadA* in dark-green leaves and its absence in pale-green leaves (not shown). The pale-green resistant plants resembled the colour of the $\Delta rbcl$ recipient used for transformation and were likely to be spontaneous spectinomycin-resistant mutants, a common problem with spectinomycin-based selection (Klaus *et al.*, 2003). Shoots were transferred to Magenta jars for rooting without further cycles of regeneration. Dark-green clones (Figure 2k) stained blue with X-Gluc (Figure 2l), indicative of the presence of the *uidA* gene, whereas the pale-green resistant shoots and recipient $\Delta rbcl$ -8a plant (Figure 2g) used for transformation were GUS-negative (Figure 2j). Leaves from dark-green pUM79 transformants contained normal levels of LS RubisCO (not shown). *Hind*III digests of pUM79 transformant DNA gave 7.0 kbp *rbcl* (Figure 4a, lanes 5 and 6) and 4.6 kbp *accD* bands (Figure 4c, lanes 5 and 6), compatible with the integration pattern of pUM79 (Figure 1b).

The absence of a 9.6 kbp *HindIII* *accD* band, diagnostic of $\Delta rbcL$ plants (Figure 4c, lanes 3 and 4) in pUM79 transformants (Figure 4c, lanes 5 and 6) indicates that the $\Delta rbcL$ plastid genomes have been replaced with a homoplasmic population of transgenic pUM79 plastid genomes. An *aadA* probe hybridized to the expected 4.1 kbp *HindIII* band in digests of DNA from pUM79 transformant clone 1 (Figure 4d, lane 5).

Discussion

We have described a precise and simple method for deleting the plastid *rbcL* gene following a single transformation step. The precision of homology-based excision is demonstrated by the presence of only five foreign bases (GGCCG) in the $\Delta rbcL$ plastid genome described here. The deletion (co-ordinates 57 424–59 317; Wakasugi *et al.*, 2001) removes the 1434 bp *rbcL* coding region, 171 bp of upstream and 289 bp of downstream sequences. This makes the mutant suitable for plastid transformation with vectors containing changes in the *rbcL* coding sequence for structure–function studies (Andrews and Whitney, 2003; Parry *et al.*, 2003; Spreitzer *et al.*, 2005) and in the regulatory region (Shiina *et al.*, 1998) to modify *rbcL* expression levels. Rescue of the mutant phenotype by the WT *rbcL* allele following *aadA*-based plastid transformation rules out mutations at other sites.

A relatively long duplication of 649 bp separated by 5.4 kbp was used because excision was barely detected between two copies of a 418 bp *psbA* 3' regulatory region spaced 1 kbp apart (Iamtham and Day, 2000). The influence of sequence composition on rates of plastid recombination is not known. High levels of sectoring in leaves of T_1 plants indicated efficient excision of the *rbcL* gene. Excision products were also detected in the T_0 generation by DNA blot analyses. This indicates that it should be possible to visualize mutant sectors in leaves of T_0 plants following vegetative propagation and additional cycles of regeneration on non-selective medium. Cytoplasmic sorting is an essential component of the method because it drives the segregation of different plastid types and leads to the isolation of homoplasmic mutants in the absence of selection. Use of site-specific recombinases is less reliant on cytoplasmic sorting because, in principle, they allow simultaneous excision of genes from multiple plastids (Kuroda and Maliga, 2003).

Deletion of native plastid genes and foreign genes in one recombination event simplifies the isolation of plastid mutants lacking foreign genes. The method has the added advantage of allowing any two regions of ptDNA to be spliced together by recombination and reduces the reliance on the proximity of convenient restriction sites to delete plastid genes. The approach also allows the deletion of multiple adjacent plastid genes. A duplication of the region

downstream of the *atpE* gene would delete the *rbcL*, *atpB* and *atpE* genes in one step. Deletion of plastid genes that do not confer a pigment-deficient phenotype, such as the *ndh* genes (Burrows *et al.*, 1998; Kofer *et al.*, 1998) would require PCR-based screening of seedlings or shoots from vegetatively propagated plants. Only regions of the plastid genome lacking essential genes can be deleted by this approach. In tobacco, essential genes include *clpP1* (Kuroda and Maliga, 2003; Shikanai *et al.*, 2001), *ycf1*, *ycf2* (Drescher *et al.*, 2000) and *accD* (Kode *et al.*, 2005) as well as plastid-encoded components of the plastid translation machinery such as *rps14* (Ahlert *et al.*, 2003) that are needed to express these genes.

Restoration of photosynthesis following *aadA*-based plastid transformation with the WT *rbcL* allele helps to maintain recombinant plastid genomes when antibiotic selection is removed in soil-grown plants. When WT plants are transformed, residual heteroplasmy in soil-grown plants can be a problem if stochastic fluctuations or selective forces lead to a reduction in transgenic plastids relative to WT plastids. One cycle of regeneration usually suffices to isolate homoplasmic transplastomic plants following mutant rescue in agreement with the results of Klaus *et al.* (2003). We were unable to use restoration of photosynthesis as the sole criterion for selecting plastid transformants (results not shown). The reasons for this failure are unclear but others have also had little success with this approach (Klaus *et al.*, 2004).

In *Chlamydomonas* (Goldschmidt-Clermont, 1991), moss (Sugiura and Sugita, 2004) and all angiosperm species in which stable plastid transformation has been reported (see below), transforming DNA integrates by homologous recombination. In addition to tobacco, stable transformation of angiosperm plastids has been reported in *Arabidopsis thaliana* (Sikdar *et al.*, 1998), carrot (Kumar *et al.*, 2004a), cauliflower (Nugent *et al.*, 2006), cotton (Kumar *et al.*, 2004b), *Lesquerella fendleri* (Skarjinskaia *et al.*, 2003), lettuce (Lelivelt *et al.*, 2005), petunia (Zubko *et al.*, 2004), potato (Sidorov *et al.*, 1999), soybean (Dufourmantel *et al.*, 2004) and tomato (Ruf *et al.*, 2001). Conservation of the homologous recombination pathway in plastids from divergent species makes homology-based excision a suitable method for manipulating ptDNA in a wide variety of plants.

Experimental procedures

Vector construction

Vectors were made by standard cloning techniques (Sambrook *et al.*, 1989). The *atpB* 5' regulatory region including the ATG initiation codon were amplified with primers *atpB-NotI-F* 5'-CCGCGCGCCGCGCAATACATCATTATTGTATAC (underlined complement of 57 423 to 57 400 bp) in 155 939 bp tobacco plastome (accession number Z00044) and *atpB-NcoI-R* CCCCATGGACATA-

ATAATAAAATAAATAAATATG (underlined positions 56 775 to 56 799 bp). The amplified product was cloned into the *EcoRV* site of pBluescript to make pBS-5'NtstpB and sequenced to ensure the absence of PCR mutations. A *uidA* coding sequence terminated by a 3' NtspbA regulatory element (Iamtham and Day, 2000) and flanked by *NcoI* and *NotI*-*SacI* sites was excised as a *NcoI*-*SacI* fragment and ligated to *NcoI*-plus *SacI*-digested pBS-5'NtstpB. The 5' NtstpB-*uidA*-3'NtspbA expression cassette was then excised with *NotI* and cloned in inverted orientation with respect to a 16SrrnBn-aadA-BnpsbC cassette (Zubko *et al.*, 2004) into the *NotI* site of pTB27-link (Zubko *et al.*, 2004) to make pUM83. In the transformation vector pUM79, the NtstpB-*uidA*-NtspbA expression cassette is replaced with a CratpA-*uidA*-NtspbA cassette, where CratpA is the 5' regulatory region present in a 680 bp *EcoRI*-*NcoI* fragment from pUC-atpX (Goldschmidt-Clermont, 1991; accession number J01399 bases 259–911). Accession numbers are AJ276677 (16SrrnBn), AJ578474 (BnpsbC), X02340 (*aadA*, modified by Goldschmidt-Clermont, 1991), U12369 (*uidA*).

Isolation of transplastomic plants

Plastid transformants were isolated following particle bombardment as described previously (Kode *et al.*, 2005). Following bombardment of WT leaves, resistant shoots were selected on RMOP medium (Svab and Maliga, 1993) containing spectinomycin dihydrochloride pentahydrate plus streptomycin sulphate (Melford Lab Ltd, Chelsworth, UK), each at 500 mg l⁻¹. Following three cycles of regeneration on RMOP medium with both antibiotics, shoots were rooted on Murashige and Skoog (MS) medium containing 200 mg l⁻¹ spectinomycin. When *ΔrbcL* leaves were used for transformation, resistant shoots were isolated on RMOP medium containing spectinomycin (500 mg l⁻¹) and transferred for rooting on MS medium with spectinomycin (200 mg l⁻¹) or MS medium lacking antibiotics after one cycle of regeneration. All solid RMOP medium was prepared with 0.7–0.8% w/v agar. Either 0.7% agar or 0.25% Phytigel (Sigma-Aldrich, Poole, UK) was used to solidify MS media for rooting. Leaf pieces were stained in X-Gluc to monitor GUS expression as described previously (Jefferson, 1987).

DNA manipulations

Total DNA was prepared using the Wizard genomic plant DNA purification kit (Promega, Madison, WI, USA). DNA blot analyses procedures, hybridization probes, PCR analyses using purified DNA or crude DNA preparations as template DNA with ReadyMix Taq containing MgCl₂ (Sigma-Aldrich, Poole, UK) were performed as described previously (Kode *et al.*, 2005). Oligonucleotides (Sigma-Aldrich, Haverhill, UK) used for primers were (F, forward; R, reverse) pt F 5'-CAACACTATCTCGACCTTGA-3' (tobacco plastid map locations 56 612–56 631), pt R 5'-GCATGAAAATACAATAGATGAATAG-3' (complement 59 797–59 821), pt F1 5'-GCAGTGGACGTTTGGATAAG-3' (59 005–59 025), aadA-F 5'-ATTCTCCGCGCTGTAGAAGT-CACC-3' and aadA-R 5'-TACATTCGCTCATCGCCAGCC-3'. PCR products and restriction fragments fractionated on 0.8–2% agarose gels were purified using the Perfectprep Gel Cleanup kit (Eppendorf AG, Hamburg, Germany). Purified PCR products and plasmids were sequenced from appropriate primers by cycle sequencing using the Big Dye terminator sequencing ready reaction kit (Applied Biosystems, Warrington, UK). DNA sequences were analysed using the Vector NTI version 9 suite of DNA analysis programs (Invitrogen, Carlsbad, CA, USA).

Microscopy

Sections were prepared as described previously (Kode *et al.*, 2005) for transmission electron microscopy using an FEI Tecnai 12 Bio-twin transmission electron microscope (FEI company, Eindhoven, The Netherlands).

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