A bacterial signal peptide is functional in plants and directs proteins to the secretory pathway

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Received 19 February 2009; Revised 27 April 2009; Accepted 30 April 2009

Abstract

The *Escherichia coli* heat-labile enterotoxin B subunit (LT-B) has been used as a model antigen for the production of plant-derived high-valued proteins in maize. LT-B with its native signal peptide (BSP) has been shown to accumulate in starch granules of transgenic maize kernels. To elucidate the targeting properties of the bacterial LT-B protein and BSP in plant systems, the subcellular localization of visual marker green fluorescent protein (GFP) fused to LT-B and various combinations of signal peptides was examined in *Arabidopsis* protoplasts and transgenic maize. Biochemical analysis indicates that the LT-B::GFP fusion proteins can assemble and fold properly retaining both the antigenicity of LT-B and the fluorescing properties of GFP. Maize kernel fractionation revealed that transgenic lines carrying BSP result in recombinant protein association with fibre and starch fractions. Confocal microscopy analysis indicates that the fusion proteins accumulate in the endomembrane system of plant cells in a signal peptide-dependent fashion. This is the first report providing evidence of the ability of a bacterial signal peptide to target proteins to the plant secretory pathway. The results provide important insights for further understanding the heterologous protein trafficking mechanisms and for developing effective strategies in molecular farming.

Key words: GFP, LT-B, recombinant proteins, secretory pathway, signal peptide, targeting, transgenic plants.

Introduction

Production of recombinant proteins such as high-value pharmaceutical or industrial products in plants has many advantages, including safety and scalability of plant production systems (Streatfield, 2007). However, achieving high levels of recombinant proteins in a heterologous system is an ongoing challenge. Transgenic protein expression is driven mostly by the selection of strong and/or tissue-specific promoters and specific targeting signals for proper compartmentalization within the cell. This allows for different levels of regulation and modifications that result in variable levels of recombinant protein expression and accumulation (Hood, 2004). The subcellular localization of a protein accounts greatly for its proper maturation and function within the cell, hence it affects the degree to which functional protein is accumulated.

Among the plant systems used for recombinant protein expression, high-yielding crops have received a lot of attention in the past decade. Maize has been demonstrated as an effective expression system for functional proteins of prokaryotic, viral, and eukaryotic origins (Murry *et al*., 1993; Hood *et al*., 1997; Kusnadi *et al*., 1998; Streatfield *et al*., 2001, 2002). The commercialization of corn-produced β-glucuronidase (Witcher *et al*., 1998), aprotinin (Zhong *et al*., 1999), and avidin (Hood *et al*., 1997) demonstrated the viability of this expression system at the commercial level. Maize has also been used for the production of edible vaccines against deadly diseases such as traveller’s diarrhea through expression of the antigenic subunit B of *E. coli* heat-labile enterotoxin (Chikwamba *et al*., 2002a, b; Streatfield *et al*., 2003). Despite concerns over using grain crops for the production of recombinant proteins, both the scientific and economic advantages of maize-based approaches are undeniable (Ramessar *et al*., 2008).

Maize seed is known for its large storage capacity and stability of proteins and starches, hence it is considered as
an ideal organ for manufacturing recombinant proteins (Ramessler et al., 2008). While there are extensive studies on maize native seed storage proteins (Shewry and Tatham, 1990; Shewry and Halford, 2002), knowledge related to heterologous protein production and storage are limited. A number of studies report the subcellular localization of the recombinant protein in the transgenic maize (Hood et al., 1997; Witcher et al., 1998) and the relationship between subcellular localization and recombinant protein yield (Streatfield et al., 2003). These reports highlighted the importance of proper compartmentalization for optimum expression and accumulation of functional protein in plants. Expression of the E. coli heat-labile enterotoxin subunit B (LT-B) gene in maize with its N-terminus native bacterial signal peptide or with a native 27 kDa γ-zein signal peptide resulted in the accumulation of LT-B in the starch granules of the transgenic kernels (Chikwamba et al., 2003). This observation has raised a question on how the bacterial protein and its signal peptide behave in the plant cellular machinery. Considerable interest has also arisen in the possible trafficking pathway of recombinant proteins in plants.

Protein translocation pathways in bacteria (Papanikou et al., 2007; Driessen and Nouwen, 2008) and plants (Raikhel and Chrispeels, 2000; Hanton et al., 2007; Hormann et al., 2007; Brown and Baker, 2008; Rojo and Denecke, 2008) have been the focus of many studies and continue to be of great interest in today’s research. In Gram-positive bacteria, secreted proteins only need to cross the plasma membrane, while in Gram-negative bacteria, secreted proteins are translocated through the plasma membrane into the periplasm and through the outer membrane. Most secreted proteins carry an N-terminal signal sequence that directs them to the inner membrane protein translocation machinery to be internalized into the membrane, translocated to the periplasm or secreted (Saier, 2006; Saier et al., 2008).

In plants, however, the sorting of proteins in the cell is somewhat more complex due to the presence of different organelles, and hence, different types of membranes. Finely orchestrated gene regulation and tissue gene expression along with subcellular compartmentalization of plant proteins allows cells to become specialized and differentiated to fulfill their role in tissue specificity, organ identity, and organism performance. Proteins carrying an N-terminal signal peptide are often directed to the secretory pathway, transported into the endoplasmic reticulum (ER) lumen, and sorted thereafter to the Golgi, cell wall, vacuole or plasma membrane, or retained in the ER. Proteins lacking a signal sequence are translated in the cytosol, and if carrying appropriate targeting signals, they are then translocated into other organelles such as the plastids (Inaba and Schnell, 2008; Paul, 2008) or mitochondria (Attardi and Schatz, 1988). Two cases have been described in which proteins are targeted to the plastid through the secretory pathway (Villarejo et al., 2005; Nanjo et al., 2006). Two native plant proteins, α-carbonic anhydrase from Arabidopsis (Villarejo et al., 2005) and rice plastidial N-glycosylated nucleotide pyrophosphatase/phosphodiesterase (Nanjo et al., 2006) were shown to localize to the plastids after being processed in the ER, describing a novel pathway for protein trafficking into the plastids. Most recently, a novel mechanism for targeting into plant mitochondria has also been reported; successful targeting of Arabidopsis thaliana mitochondrial-targeting signal 1 (MITS1) protein to mitochondria is influenced by an N-terminal extension serving as a targeting peptide as well as by domains in the full-length protein (Chatre et al., 2009). The MITS1 N-terminal extension was shown to contain three regions that coordinate the mitochondrial targeting signal, including a cryptic signal for protein targeting to the secretory pathway (Chatre et al., 2009).

Although much has been learned about the translocation machineries of bacteria and plants, less is known about how plants utilize their molecular translocation machineries to sort bacterial proteins when expressed in transgenic plants. LT-B has a native signal peptide that directs the protein for secretion in bacteria. In E. coli, LT-B is exported to the periplasm in a Sec-dependent fashion via the general secretory pathway. Once the signal peptide is cleaved, the protein assembles into a functional pentamer and associates with the A subunit (LT-A) to form the holotoxin before being secreted from the cell via a type II protein secretion pathway (Tauschek et al., 2002). LT-B has been expressed in several plant species such as potato (Mason et al., 1998; Tacket et al., 1998), tobacco (Kang et al., 2003), maize (Chikwamba et al., 2002b; Streatfield et al., 2003) and soybean (Moravec et al., 2007). Although several of these studies describe the targeting of LT-B to different cellular compartments by the inclusion of targeting signals within their constructs (Streatfield et al., 2003), only two studies show the detailed subcellular localization of LT-B in the transgenic material (Chikwamba et al., 2003; Moravec et al., 2007). Streatfield and colleagues (Streatfield et al., 2003), for example, used the signal sequences from barley α-amylase and aleurin to target LT-B to the cell surface and vacuole, respectively, and the maize granule-bound glycoprotein synthase to target LT-B to the plastids. In the Chikwamba work (Chikwamba et al., 2003) the functional LT-B was found in the starch granules of transgenic maize kernels expressing the LT-B with its native bacterial signal peptide. In the Moravec work (Moravec et al., 2007) the replacement of bacterial signal peptide with an Arabidopsis basic chitinase signal peptide at the N-terminus of LT-B resulted in the localization of LT-B in protein bodies of transgenic soybean seeds.

The focus of this study is to examine the subcellular trafficking of the bacterial LT-B protein and its native signal peptide in plant systems using the green fluorescent protein (GFP) as a reporter. Using the functional LT-B::GFP fusion proteins it is shown that the LT-B signal peptide, not LT-B protein itself, can direct cargo proteins to the secretory pathway in Arabidopsis thaliana and maize. In addition, in maize seed, the bacterial signal peptide leads to a strong association of the cargo proteins with the starch fraction, although most of the soluble fusion protein is found in the fibre fraction. The results provide insights for further
understanding the processing of a bacterial protein in the plant cells and future design of a high level production system for recombinant proteins in plants.

Materials and methods

DNA constructs

A schematic representation of the constructs used in this study is presented in Fig. 1. The enhanced green fluorescence protein (EGFP) sequence in pLM01, pLM02, pLM03, pLM08, and pLM09 was cloned from p27zn-signal (Shepherd and Scott, 2009) using standard molecular biology techniques for restriction enzyme-based cloning. Construct pTH210 containing the CaMV 35S promoter (P35S), tobacco etch virus translational enhancer (TEV) and LT-B was used as a cloning vector (Mason et al., 1998) for some of the constructs. For the generation of plasmids pLM01, pLM02, and pLM03 the EGFP sequence was cloned into pTH210 at the NcoI–SacI sites, SacI–SacI sites, and KpnI–SacI sites, respectively. Plasmid pLM01 was used as a backbone for the generation of pLM08 and pLM09. Plasmid pRC5, a pUC19-based vector carrying the maize 27 kDa γ-zein promoter (Pγzein), TEV, the maize 27 kDa γ-zein signal peptide (ZSP) fused to LT-B and the soybean vegetative storage protein terminator (Tvsp), was used as a source of LT-B fused to the maize 27 kDa γ-zein signal peptide (R Chikwamba, unpublished results). The ZSP-LT-B fragment was amplified by the polymerase chain reaction (PCR) method and the digested product was cloned into the NcoI–BstXI site of pLM01. An NcoI–EcoRI fragment of pLM08 was inserted into the NcoI–EcoRI backbone of pLM01 to generate pLM09. A simple alanine-glycine linker (AG linker) consisting of six amino acids with three AG repeats (AGAGAG) was added between LT-B and GFP using oligonucleotide extensions in the PCR primers in plasmids pLM03, pLM08, and pLM09. Expression cassettes (Fig. 1) were cloned and recombined into a Gateway version of pTF101.1 (Paz et al., 2004), pTF101.1gw1, for Agrobacterium-mediated transformation of Arabidopsis thaliana. Plasmid pRC4 (Chikwamba et al., 2002b) was used as a donor of the Pyzein, and derived plasmids.

The nuclear marker VirD2::RFP was a kind gift of Dr Stanton Gelvin. The ER marker plasmid designated here as ER cherry (Nelson et al., 2007) was obtained from the Arabidopsis Biological Resource Center (ABRC Stock No. CD3-959, http://arabidopsis.org).

Stable transformation of maize

Maize transformation was carried out at the Center for Plant Transformation at Iowa State University as described previously (Frame et al., 2000). Briefly, DNA constructs (Fig. 1) were co-bombarded with a selectable marker gene that confers resistance to the herbicide bialaphos. Herbicide-resistant calli were analysed using PCR for presence of the GFP and LT-B genes. For constructs driven by P35S, the calli of transgenic events were tested for GFP fluorescence and imaged as described in the ‘Fluorescence imaging and laser scanning confocal microscopy’ section of the Materials and methods. Transgenic calli were regenerated and plants brought to maturity in the greenhouse.

Protein extraction from transgenic tissue

Endosperm powder samples were collected from transgenic kernels using a hand-held drill as described by Sangtong et al. (2002). Plant materials were incubated with the following protein extraction buffer at a rate of 10 μl buffer

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<td>P314</td>
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Fig. 1. Gene cassettes, plasmids, and transgenic lines for studying localization of LT-B using GFP as a reporter. Gene cassettes A–E were used in transient assays using Arabidopsis leaf and root protoplasts, and in stable transformation of maize callus and endosperm tissues. Constitutive expression in Arabidopsis protoplasts and in maize callus was driven by the double CaMV 35S promoter (P35S promoter). The maize 27 kDa γ-zein promoter (Pγzein promoter) was used to drive expression of gene cassettes A–E in maize endosperm. The number of independent transgenic events for each line is presented in parenthesis. TEV, tobacco etch virus translational enhancer leader sequence; EGFP, enhanced green fluorescent protein; Tvsp, the soybean vegetative storage protein terminator; BSP, the LT-B bacterial signal peptide; LT-B, the B subunit of E. coli heat-labile enterotoxin; ZSP, 27 kDa γ-zein signal peptide; AG linker, alanine-glycine linker; na, not available.
mg⁻¹ maize powder: 25 mM sodium phosphate (pH 6.6), 100 mM NaCl, 0.1% Triton X-100 (v/v), 1 mM EDTA, 10 μg ml⁻¹ leupeptin, and 0.1 mM serine protease inhibitor Perfabloc SC (Fluka), for 2 h at 37 °C. Total aqueous extractable protein (TAEP) was determined using the Bradford assay (Bradford, 1976).

**LT-B detection by Gm₁ capture ELISA**

Quantification of LT-B in the samples was carried on using a modification of the monosialoganglioside-dependent enzyme-linked immunosorbent assay (ELISA) described previously (Chikwamba et al., 2003). The described protocol was modified as follows: monosialoganglioside, Gm₁, from bovine brain (G7641, Sigma, St Louis, MO, USA) was used at a 10 μg ml⁻¹ concentration, 50 μl per well. Streptavidin–horseradish peroxidase conjugate (554066, BD Biosciences, San Jose, CA, USA) was used at a dilution of 1:1000 in 1% dry milk (DM) (w/v) in phosphate buffered saline [PBS: 0.01 M Na₂HPO₄, 0.003 M KH₂PO₄, 0.1 M NaCl (pH 7.2)]. Horseradish peroxidase substrate [ABTS; 0.5 mM 2,2-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid) diammonium salt, 0.1 M citric acid, pH 4.35] was activated prior to use, by adding 5.5 μl 30% H₂O₂ to 5.5 ml ABTS solution. Activated horseradish peroxidase substrate was added to the plate and incubated in the dark at room temperature for 30 min. Absorbance was measured spectrophotometrically at 405 nm at the end of the reaction. Sample wells were blanked against non-transgenic maize protein extracts and all measurements were performed in duplicate. Raw ELISA data was converted to per cent LT-B of total soluble protein by reference to an ELISA standard curve constructed using purified bacterial LT-B (kindly provided by Dr John Clements, Tulane University, LA, USA).

**Western blotting**

An aliquot containing 50–100 μg of total aqueous extractable protein (TAEP) from maize kernels was boiled for 5 min and loaded onto a 15% polyacrylamide SDS-PAGE (Laemmli, 1970). The separated proteins were transferred to a 0.45 μm Nylon Filter (146506, Spectrum Consultants Laboratory, Inc., Newberg, OR, USA) over-night in the cold room (9–11 °C). Protein G beads (IP50-1KT, Sigma, St Louis, MO, USA) were used to recover the immunoprecipitation (IP) complex, following the supplier’s recommendations. Ten μl of recovered IP complex were analysed by Western blotting using goat anti-GFP antibody (Cat no. 600-101-215, Rockland Immunocchemicals, Inc., Gilbertsville, PA, USA) as described in the ‘Western blotting’ section of the Materials and methods.

**Thermolysin treatment of starch samples**

Thermolysin (3097-ZN, R&D System, Inc., Minneapolis, MN, USA) was diluted to 10 μg ml⁻¹ with 5 mM CaCl₂ solution. Ten milligrams of starch were incubated with 100 μl thermolysin solution at 37 °C for 2 h. The reactions were terminated by the addition of EDTA to a final concentration of 20 mM. Samples were subsequently washed five times with 1 ml distilled water. After the last wash, 100 μl 1× SDS sample buffer was added to each sample and boiled for 5 min. Twenty μl were loaded onto an SDS-PAGE gel for
Western blot analysis, as described in the ‘Western blotting’ section of the Materials and methods.

*Arabidopsis thaliana* mesophyll and root culture protoplast isolation for transient transformation assays

*Arabidopsis thaliana* (ecotype Columbia) seeds were vernalized in water at 4 °C for 48 h before sowing in prewetted LC1 Sunshine Mix. Flats were placed in a growth chamber at 21 °C and 16 h photoperiod covered with humidomes for 2–3 d. One to two weeks after germination, plants were thinned and transplanted to individual pots for further growth. Isolation of mesophyll protoplasts was carried on as described by Sheen (2001).

*Arabidopsis* root protoplasts were isolated from a root culture that was maintained on Lindmaier–Skog media (LSP003, Caisson Laboratories, North Logan, UT, USA) supplemented with 20 g l⁻¹ sucrose, 1 mg l⁻¹ naphthalene-acetic acid (NAA), and 0.05 mg l⁻¹ kinetin (Contesto et al., 2005), kindly provided by Dr D Bassham at Iowa State University. For the isolation of root culture protoplasts, 10–20 ml of root cell culture was pelleted in a 50 ml Falcon tube by centrifuging for 4 min at 50 g, or resting on the bench. The supernatant was removed and replaced by 40 ml of fresh enzyme solution [0.5× artificial sea water (ASW, 1.7 M NaCl, 9.4 mM MgSO₄, 3.4 mM CaCl₂, 5 mM MES, 3.45 mM KCl, 8.35 mM MgCl₂·6H₂O, 0.875 mM NaHCO₃, pH 6.0), 0.3 M mannitol, 0.31% Cellulase Onozuka R10, 0.15% Macerozyme R10, pH 5.7, filter sterilized]. Cells were transferred to a deep Petri dish and were vacuum infiltrated for 5 min. The Petri plate was covered with aluminum foil and placed on an orbital shaker for 3.5 h at 50 rpm. The digested cell solution was filtered through a 40 lµm mesh into a 50 ml conical Falcon tube, and the plate was washed once with cold W5 solution (154 mM NaCl, 125 mM CaCl₂, 5 mM KCl and 2 mM MES, pH 5.7, filter sterilized). Cells were spun at 50 g for 4 min. The supernatant was removed and cells were washed twice with W5. The cells were carefully resuspended in fresh W5 and incubated on ice for at least 30 min prior to transformation with polyethylene glycol (PEG). Leaf and root protoplasts were transformed using PEG as described in Sheen (2001).

Sample preparation for microscopy

Transformed mesophyll and root culture protoplasts of *Arabidopsis thaliana* were washed and resuspended in fresh W5 media. Thirty µl of protoplast solution were placed directly on the microscope slides for visualization. Transgenic maize callus and developing endosperm were mounted directly onto microscope slides using Vecta Shield mounting media with propidium iodide (H1200, Vector Laboratories, Burlingame, CA, USA).

Fluorescence imaging and laser scanning confocal microscopy

*Arabidopsis* protoplasts, maize callus, and maize endosperm samples were visualized using a Leica TCS/NT (Leica Microsystems Inc., Exton, PA, USA) laser scanning confocal microscope equipped with Argon (488 nm) and Krypton (568 nm) lasers and a double dichroic DD488/568 filter. Green fluorescence was detected using a combination of RSP580 and BP525/25 filters under wavelengths between 500 and 550 nm. Red fluorescence was detected using a LP590 filter. During scanning, pinhole was maintained at 1 (airy units) for all images. Images were processed and analysed using ImageJ software (Abramoff et al., 2004).

Ears from self-pollinated transgenic maize plants harvested at 10–14 d after pollination were visualized and imaged using an Olympus SHZ10 stereo scope (Leeds Precision Instruments, Inc., Minneapolis, MN, USA) coupled to a SPOT RT colour CCD camera (Diagnostic Instrument Inc., Sterling Heights, MI). Images were taken under bright field or using a band pass exciter at 460–490 nm with emission filter 510–550 nm (for GFP detection), and acquired using SPOT Advanced software.

Hydropathy plots, secondary structure, and subcellular localization prediction

Hydropathy scores for signal peptide sequences of BSP (signal peptide of the B subunit of *E. coli* heat-labile enterotoxin; gi:88687064), ZSP (signal peptide of the maize 27 kDa γ-ztein; gi:16305109), and BiPSP (signal peptide of the maize luminal binding protein; gi:1575128) were obtained based on Kyte and Doolittle (1982) using ProtScale prediction software (Gasteiger et al., 2005). Hydrophobicity plots were generated in Excel. Secondary structure for the aforementioned signal peptides were predicted using PSIPred (Jones, 1999; McGuffin et al., 2000). Prediction of subcellular localization was done using TargetP 1.1 (Emanuelsson et al., 2000; Nielsen et al., 1997) and iPSort (Bannai et al., 2002) software.

Results

Protein fusions of heat-labile enterotoxin subunit B (LT-B) and the green fluorescent protein (GFP) assemble and fold properly in plant cells

Constructs listed in Fig. 1 were made to investigate the subcellular targeting properties of the LT-B signal peptide and the LT-B protein using GFP as a visual marker. The first two constructs have GFP without a signal peptide (Fig. 1A, control), or fused to the bacterial signal peptide (BSP) of LT-B (Fig. 1B). The last three constructs have GFP fused at the C-terminus of the full-length LT-B protein with its native BSP (Fig. 1C), a maize 27 kDa γ-ztein signal peptide (ZSP) (Fig. 1D), or no signal peptide (Fig. 1E).

To facilitate rapid construct verification and protein localization in different plants, each construct set was made with two different promoters. A constitutive CaMV 3SS promoter was used for transient analysis in *Arabidopsis* protoplasts and stable analysis in transgenic maize callus culture, while a maize seed-specific 27 kDa γ-ztein promoter was used for endosperm analysis in transgenic kernels. For
each construct, a transgenic line identification was designated and multiple independent transgenic events were generated (Fig. 1). All transgenic maize lines were confirmed by gene-specific PCR analysis. The transgene insertion copy numbers were estimated by Southern blot analysis (data not shown). Strong transgene expressing events were chosen for further localization and biochemical analyses.

To examine whether LT-B and GFP retain their functional properties in the fusions described, experiments were carried out to assess their correct folding and assembly in maize. In stably transformed callus, green fluorescence could easily be distinguished when the fusion constructs were present (data not shown). Similarly, it was possible to detect green fluorescent transgenic maize seeds (Fig. 2A) confirming that GFP, a molecule capable of folding and fluorescing without exogenous substrates or cofactors (Chalfie et al., 1994) is active and correctly folded.

A monosialoganglioside (GM₁) capture Enzyme Linked Immunosorbent Assay (ELISA) was used to determine whether the LT-B protein assembles into functional pentamers when fused to GFP protein. To be biologically functional, the 11.7 kDa LT-B protein needs to be correctly assembled into pentameric form, which can then bind to monosialoganglioside specifically (Spangler, 1992). Figure 2B shows that multiple events of transgenic maize lines P310 and P311 accumulate pentameric LT-B at various levels.

These results combined indicate that both LT-B and GFP were able to retain their properties when fused; LT-B being in a pentameric form remains recognizable by its receptor, the G₉₁ receptor, and GFP retains its characteristic of self-catalysis for folding and fluorescing.

Western blot analysis using anti-GFP and anti-LT-B antibodies on total aqueous extractable protein (TAEP) (Fig. 2C, D) were conducted to further confirm the presence of the fusion proteins in transgenic maize kernels. As shown in Fig. 2C, using anti-GFP antibody it was possible to detect bands around 27 kDa (arrowheads) in transgenic lines P309 (P₇₇-GFP control) and P315 (P₇₇-BSP-GFP) as predicted. In P315, two bands with similar molecular weights probably represent populations of GFP with and without cleaved signal peptide. On the other hand, in transgenic maize lines P308c (P₃₅S-BSP-LT-B::GFP), P310-26 and -32 (P₇₇-BSP-LT-B::GFP), and P311 (P₇₇-ZSP-LT-B::GFP), the analysis also detects a number
of bands that cross-reacted to anti-GFP antibody. One high molecular weight band estimated around 39 kDa (asterisks in Fig. 2C) is possibly the LT-B (11.7 kDa) and GFP (26.9 kDa) fusion product. Bands at the lower molecular weights (dots in Fig. 2C) cross-reacting to the antibody suggest possible proteolytic degradation of the fusion protein in these transgenic lines. No signal was observed for B73, a non-transgenic maize inbred line.

The anti-LT-B antibody (Fig. 2D) cross-reacted with bands around 39 kDa in P308c and P310 (asterisks in Fig. 2D), which are probably the LT-B::GFP fusion proteins. In P308c callus, a lower molecular weight band around 11 kDa was also detected (closed diamond in Fig. 2D). Although the size of this band is similar to the monomeric form of LT-B (open diamonds in Fig. 2D) as shown in bacterial LT-B standard and P77, a control transgenic maize line carrying LT-B with its own BSP (Chikwamba et al., 2002b), current evidence is not enough to show that it represents a processed LT-B monomer. No LT-B cross-reacting bands are seen in constructs carrying GFP alone (P309 and P315), as expected.

Figure 2E shows an anti-GFP Western blot on immunoprecipitated transgenic maize kernel samples using anti-LT-B antibody. A reacting band was detected around the 39 kDa position in maize lines P310 and P311, but no band could be seen in two control maize lines P309 (Pγzein-GFP) and P77 (Pγzein-LT-B, Chikwamba et al., 2002b) carrying GFP or LT-B alone, respectively. This result confirms that the ~39 kDa bands observed in Fig. 2C are, in fact, fused to LT-B, as predicted. Figure 2E included a commercially EGFP standard (Cat no. 4999-100; Biovision, Mountain View, CA, USA), which shows more than one cross-reacting band when probed using goat anti-GFP antibody (arrow). This phenomenon was observed in other Western blots when using the anti-GFP antibody in our work. It is possible that the multiple GFP bands in commercial EGFP standard are the result of incomplete protein denaturation during boiling before loading.

**LT-B::GFP fusion proteins are detected in starch and fibre fractions of transgenic maize kernels**

Starch and fibre fractions were analysed for functional GFP and LT-B content in transgenic lines P310 (Pγzein-BSP-LT-B::GFP) and P311 (Pγzein-ZSP-LT-B::GFP). The TAE of all samples was obtained by two different extraction methods: simple vortex at room temperature (25 °C) and shaker-incubation at 37 °C for 2 h. These two methods of protein extraction were chosen to investigate the stability of the fusion protein by the previously described method of LT-B extraction (2 h/37 °C, Chikwamba et al., 2002b, 2003) compared to a more conservative extraction protocol (vortex/RT). As shown in Fig. 3A, the percentage of LT-B of TAE obtained from 2 h/37 °C incubation was generally higher than that from vortex/RT extraction, suggesting that a better soluble protein recovery was achieved when using a method with higher temperature and a longer incubation time. Figure 3A also indicates that LT-B is not only associated with the starch fraction as previously reported, but also associated, at a higher percentage, to the fibre fraction obtained using the small-scale fractionation protocol. This association was also observed for the LT-B transgenic maize line P77 (Pγzein-BSP-LT-B; data not shown), the original line used in the previous work to demonstrate the LT-B association to the starch granules of transgenic maize kernels (Chikwamba et al., 2003). The observed LT-B/fibre association was not reported in the previous work due to the fact that protein recovery from non-soluble and non-starch fractions was not determined.

Western analysis using the anti-GFP antibody on proteins extracted from the aqueous phase and the insoluble pellet phase of each fraction (Fig. 3B, C) were performed to confirm the presence of the fusion proteins in starch and fibre. Interestingly, strong fusion protein bands were
observed in P310 and P311 in both starch pellets (asterisks in Fig. 3B) and fibre pellets (asterisks in Fig. 3C). Weak but detectable fusion protein bands can be seen in the fibre TAEP of P310 and P311 (asterisks in Fig. 3C). However, while it was detectable by Gm1 capture ELISA specific for LT-B detection, no protein bands were observed from the starch TAEP for both P310 and P311 (Fig. 3B). As for the control construct, the GFP protein from line P315 (Pγzein-BSP-GFP) behaved similar to P310 (Pγzein-BSP-LT-B::GFP) in the Western analysis. No detectable GFP band was seen for P309 (Pγzein-GFP), which carries the GFP with no signal peptide.

Thermolysin (EC 3.4.24.27) treatment based on the published protocols (Chikwamba et al., 2003) was used to verify whether the detection of the GFP and its fusion protein in starch pellets was due to internalization in starch grains of the insoluble phase, as observed previously for LT-B alone (Chikwamba et al., 2003). Any polypeptides within the starch granules should not be susceptible to hydrolysis upon treatment of intact granules with exogenous proteases such as Thermolysin (Mu-Forster and Wasserman, 1998). Starch pellets from three transgenic lines, two carrying GFP constructs (P309 and P315) and one carrying the LT-B::GFP fusion construct (P310), were treated with Thermolysin. To monitor the Thermolysin effectiveness, maize waxy protein (Mu-Forster et al., 1996) was used as an internal control for starch-bound proteins and γ-zeins (Mu-Forster and Wasserman, 1998) were used as markers for starch granular external protein contamination. The fibre fraction of P315 was used as a treatment control. Both treated and untreated samples were subjected to Western blots that cross-reacted with anti-GFP, anti-waxy, and anti-γ-zein antibodies. If the proteins are internalized in the starch granules, the reacting band should remain at the same intensity in Thermolysin-treated and untreated samples.

As can be seen in Fig. 4A, the GFP band in the Thermolysin-treated control sample P315F (fibre fraction) was reduced. However, the P310 fusion protein band (asterisks in Fig. 4A) and P315 GFP band (arrowheads in Fig. 4A) in the Thermolysin-treated and untreated starch samples remained at the same intensity, suggesting that these proteins were not sensitive to Thermolysin treatment, and most likely are protected by the starch granule membrane.

Because the waxy protein is strongly associated with the starch granule, Thermolysin treatment did not reduce its protein band intensity in all samples (block arrow in Fig. 4B). On the other hand, the band intensity of the 16 kDa and 27 kDa γ-zeins (open block arrow in Fig. 4C), known to accumulate in protein bodies and not inside starch granules, was notably reduced upon treatment with the protease (Fig. 4C).

**The bacterial signal peptide of LT-B is sufficient for localization of GFP and LT-B::GFP fusions to the secretory system of Arabidopsis and maize**

Confocal microscopy was used for localization studies in transiently transformed *Arabidopsis* protoplasts, and stably transformed maize callus and endosperm tissue using the constructs described in Fig. 1. The objective of such experiments was to establish the subcellular targeting properties of the bacterial signal peptide of LT-B and the LT-B protein itself. Figure 5 summarizes the results.

Figure 5A is the GFP control construct in which the gfp gene is under the control of either P35S or Pγzein promotors with no signal peptide. The transient expression of GFP in *Arabidopsis* leaf (Fig. 5Aa) and root (Fig. 5Ab) protoplasts, and stable expression of GFP in maize callus (Fig. 5Ac) and endosperm (Fig. 5Ad) resulted in localization of GFP in the cytoplasm and nucleus; this observation has been reported and reviewed extensively (Hanson and Kohler, 2001; Berg et al., 2008).

The fusion of GFP to the C-terminus of the LT-B signal peptide BSP (Fig. 5B) resulted in localization of the GFP signal in the endoplasmic reticulum (ER) in all types of transformed cells tested in this study (Fig. 5Be–h).
To understand whether the mature LT-B protein also plays a role in subcellular localization in plant cells, the LT-B::GFP fusion construct linked with either bacterial BSP (Fig. 5C), plant ZSP (Fig. 5D) or no signal peptide (Fig. 5E) was assessed in various transiently or stably transformed plant cells. The results show that the GFP signals were accumulated in the secretory systems for either BSP (Fig. 5Ci–l) or ZSP (Fig. 5Dm–p) led LT-B::GFP fusion proteins in both Arabidopsis and maize, similar to the observations presented Fig. 5Be–h. On the other hand, when the signal peptide (Fig. 5E) was removed, the GFP signal returned to the cytosol and nucleus, and no signal was observed in the secretory pathway (Fig. 5Eq–s).

Figure 6 shows the co-localization experiments for the confirmation of subcellular localization using a known ER marker (Nelson et al., 2007). This marker (ER cherry)
contains a red fluorescent protein fused to the signal peptide of the Arabidopsis thaliana wall-associated kinase 2 (He et al., 1999) and an HDEL motif for retention in the ER (Munro and Pelham, 1987). Arabidopsis root protoplasts were co-transformed with the ER cherry marker construct and the various constructs listed in Fig. 1. The results show that the ER-targeted RFP signals co-localized with the GFP signals from the constructs that contained either BSP (Fig. 6B, C) or ZSP (Fig. 6D). By contrast, GFP signals from constructs carrying GFP alone (Fig. 6A) or LT-B::GFP fusion (Fig. 6E) with no signal peptide were constantly detected in separate subcellular compartments than ER-targeted RFP signal.

**Discussion**

The aim of this study was to determine the targeting properties of the B-subunit of E. coli heat-labile enterotoxin...
(LT-B) and its signal peptide (BSP) in plant cells using GFP as a visual marker. The main findings of this study are as follows: (i) BSP directs LT-B and GFP to the secretory pathway of plant cells; this is the first report of a bacterial signal peptide carrying this function in plants; (ii) LT-B protein alone has no obvious targeting properties; (iii) GFP fused to BSP, and LT-B::GFP fusions carrying BSP or a maize zein signal peptide (ZSP) can be found in starch granules of transgenic maize kernels, but are mostly strongly associated with the fibre fraction; and (iv) it is possible to generate successful functional translational fusions of LT-B (11.7 kDa) with GFP (−27 kDa); this property of LT-B has great potential applications in the biotechnology industry. Our results indicate that subcellular localization of LT-B in plant cells is determined by the presence (or absence) of a signal sequence such as BSP or ZSP. The findings also indicate that BSP plays a targeting role in plants similar to its role in bacteria, directing proteins (GFP and LT-B) to the secretory pathway. Finally, the findings also indicate that LT-B can be fused to proteins as large as 27 kDa and maintain functionality, a property highly desirable for the potential use of LT-B as a carrier molecule.

BSP acts as a functional signal peptide in plants

Signal peptides control the entry of all proteins to the prokaryotic and eukaryotic secretory pathways (Nielsen et al., 1997). The predicted secondary structure (PSIPred; Jones, 1999; McGuffin et al., 2000) of the signal peptide of LT-B (BSP) was compared with two signal peptides from maize proteins, 27 kDa γ-zein and luminal binding protein (Fig. 7A). It can be seen that all three signal peptides have common coil–helix–coil motifs, and that all three of them follow the (−3, −1) rule: the residues at positions −3 and −1 from the cleavage site must be small and neutral (Nielsen et al., 1997). Figure 7B also shows the hydropathy plot for these signal peptides. A clear hydrophobic region comprising residues 7 to 17 is consistent with the predicted helical motif. Furthermore, subcellular localization prediction softwares TargetP 1.1 (Emanuelsson et al., 2000) and iPSort (Bannai et al., 2002) identified the sequences used in constructs B–D (Fig. 1) as having a signal peptide that would target the fusion proteins to the secretory pathway. By contrast, constructs lacking a signal peptide were predicted to accumulate in the cytosol.

In E. coli enterotoxigenic strains, BSP is the signal peptide that directs LT-B protein to be translocated to the periplasm where the BSP is cleaved off, and the LT-B protein assembles into functional pentamers before being secreted in a folded state (Tauschek et al., 2002). The results presented in Figs 5 and 6 show that BSP plays the role of a signal peptide in plants, and results in the localization of the fusion proteins in the secretory pathway of Arabidopsis and maize. It is also clear that the mature LT-B protein has no targeting ability for the secretory system, and that the signal peptide alone, is sufficient for targeting cargo proteins to the ER. All the constructs used lacked an ER retention motif, which raises interesting questions regarding the ER retention observed. Proteins carrying a signal peptide are usually transported further downstream in the secretory pathway unless they carry an ER retention motif, interact strongly with ER resident proteins, or fail to fold properly (Pagny et al., 1999). Therefore, in the case of our constructs carrying a BSP or ZSP, they were expected to be secreted from the cell. It has been reported that secreted forms of GFP reaching the apoplast showed none to low fluorescence levels due to the effect of pH on GFP folding (Batoko et al., 2000; Zheng et al., 2004). Hence, it cannot be ruled out that some of the GFP from BSP- or ZSP-carrying constructs were secreted but not visualized due to unfavourable experimental conditions (e.g. different pH). At the same time, it is also possible that the LT-B::GFP fusions, in particular, might require strong interaction with ER resident chaperones and protein isomerases for proper folding and assembly, thereby resulting in accumulation in the ER lumen, as has been shown for other proteins (Saito et al., 2009).

The observation of the ER localization properties of the LT-B signal peptide may explain why a great proportion of LT-B can be detected in the fibre fraction during kernel fractionation. The fibre fraction generated in the small-scale starch preparation process contains most maize cell components except for the starch, after removal of germ and pericarp.
by manual excision. The results do not explain, however, why a fraction of GFP and LT-B::GFP proteins fused to BSP was also found in the starch granules of maize seeds. Starch granules are inside the amyloplasts, specialized starch storage plastids found in cells of the maize endosperm. Proteins targeted to the ER have an N-terminal signal peptide that guides its insertion and translocation into the ER lumen, where the signal peptide is cleaved (Vitale and Boston, 2008). The ER is the point of entry for the secretory pathway, which includes ER, Golgi, plasma membrane, vacuole, cell wall or any body derived from any of these (Robinson et al., 2007). With few reported exceptions, nuclear proteins targeted to the plastids are translated in free ribosomes in the cytosol, and are then translocated using the organelle’s specialized import machinery in a transit peptide-dependent fashion.

The BSP-dependent dual localization observed in this work has some similarity to plant signal peptide-dependent dual localization reported for rice amylases rAmy3 (Chen et al., 2004) and rAmy1-1 (Asatsuma et al., 2005) which accumulate in plastids and cell wall/extracellular space in plant cells. It remains to be determined whether the dual localization property of LT-B involves one or more trafficking routes, reaching the plastids via an ER-independent pathway. Recently, two groups described a novel pathway for protein trafficking into the plastids via the secretory pathway. Villarejo and colleagues (2005) showed that α carbonic anhydrase (α-CA) accumulates in the chloroplast stroma after being processed within the secretory system (Villarejo et al., 2005). In a different study it was shown that nucleotide pyrophosphatase/phosphodiesterase (NPP1) of rice and barley is glycosylated in the secretory pathway (Nanjo et al., 2006) before final accumulation in the plastids through a vesicular transport pathway from the ER and Golgi (Nanjo et al., 2006).

It is also possible that the default localization of the BSP-driven proteins is the ER, as was observed in Arabidopsis and maize, but additional fates occur in maize endosperm upon saturation of the secretory system. Overexpression of glycmin in soybean resulted in increased formation of ER-derived protein bodies carrying intermediates destined for the protein storage vacuole (Kinney et al., 2001). It is proposed that this response is most likely due to the formation of insoluble protein aggregates in the ER (Kinney et al., 2001). Because a large proportion of the fusion proteins in this study are found in the insoluble fraction of the endosperm, it cannot be ruled out that the dual localization can be a result of overexpression.

Unexpected patterns of recombinant protein deposition in endosperm have been reported for other species such as wheat (Arcalis et al., 2004) and rice (Drakakaki et al., 2006). In wheat, KDEL-tagged recombinant serum albumin was shown to accumulate in the ER lumen in leaves but was detected in prolamin aggregates inside vacuoles in the endosperm, as was observed with recombinant phytase targeted for secretion (Arcalis et al., 2004). In the same wheat study, recombinant legumin targeted to the vacuole resulted in the accumulation of the protein in globulin inclusion bodies around the prolamin bodies (Arcalis et al., 2004). Further effects of tissue type on recombinant protein subcellular fate was studied in rice by the expression in leaves and endosperm of recombinant phytase targeted for secretion. While, in leaves the phytase was successfully secreted, in endosperm it was retained in the endoplasmic reticulum-derived protein bodies and protein storage vacuoles (Drakakaki et al., 2006).

Because fractionation experiments were conducted specifically with transgenic maize kernels and not with Arabidopsis tissue, at this time it is difficult to establish if the observations presented for BSP-driven proteins are restricted to maize endosperm. Fractionation experiments combined with GFP localization suggest that the majority of the protein fusion population carrying a signal peptide is targeted to the ER. Using confocal microscopy, it was not possible to detect any GFP signal in plastids of either Arabidopsis or maize, but biochemical evidence suggests that a small fraction of the population was indeed associated with the starch fraction in maize kernels. In maize, the native γ-zein proteins are retained in the ER-lumen and accumulate in ER-derived protein bodies as a result of protein–protein interactions with other zeins and chaperones (Kim et al., 2002). Unlike the zein protein, the LT-B::GFP mature protein may or may not have the ability to interact with other zeins and ER resident proteins to be retained effectively in the ER or its derived systems.

BSP driven GFP and LT-B::GFP fusions associate with fibre and starch

Small-scale (5–10 kernels) starch isolation of maize kernels using a modification of the previously described protocol (Chikwamba et al., 2003) gives two main fractions: starch and fibre. As in other wet milling fractionation procedures (Johnson, 2000), this protocol allows separation and recovery of starch, fibre, germ, steep water, and pericarp. For practical purposes in this study, pericarp, embryo, and steep water, were discarded, and the endosperm was used for further separation of starch and fibre. While the starch fraction derives mostly from the amyloplasts, less is known about the cellular components of the fibre fraction.

The results presented in Fig. 3 indicate that fusions carrying BSP or ZSP are strongly associated with the fibre fraction of transgenic maize kernels, and to a lesser degree, with the starch fraction as well. While accumulation of LT-B in starch granules had been reported before (Chikwamba et al., 2003) the strong association with the fibre fraction is a novel observation, consistent with the ER-localization determined by confocal microscopy (Figs 5, 6). The strong association of LT-B with the fibre fraction may provide unique opportunities for effective downstream processing of a potential edible vaccine product, adding utility to the low value fibre fraction that results from corn fractionation (L Johnson, personal communication). The low association of LT-B with starch also presents potential advantages for the industry, as it might allow for utilization of the starch fraction for traditional applications such as biorefinery and non-food, non-feed processing.
LT-B can be used as a carrier molecule for peptide with size as large as 27 kDa

Expression of recombinant proteins in plants continues to be an area of great interest in the pharmaceutical, industrial, and vaccine industries. In this study, GFP is used as a visual marker to study the subcellular localization of the LT-B protein and its bacterial signal peptide. The data indicate that the C-terminus fusion of LT-B with GFP (using a simple alanine-glycine linker) results in retention of properties and assembly for both proteins. Because the pentameric LT-B protein is a potent antigen (Nashar et al., 1998) and binds specifically to the G_{H1} receptor of the epithelial cell surface, the results presented here suggest that LT-B has the potential to be used as a carrier molecule to deliver proteins larger than itself.

The use of LT-B as a carrier molecule for vaccination purposes has been extensively studied in bacterial systems (Buddenborg et al., 2008; Spangler, 1992). However, reports of plant-derived LT-B fusions are limited. Two studies report the fusion of small molecules (5–6 kDa) to the carboxy terminus of LT-B in Arabidopsis (Rigano et al., 2004) and in tobacco (Rosales-Mendoza et al., 2009). In both of these cases, the peptides used in the fusion represent about 1/5 of the size of the GFP protein (27 kDa) used in this study. Our work presented here is an important contribution to the vaccine research, because it is demonstrated that the LT-B can be fused with a protein as large as GFP and both proteins can retain their native conformations and functionalities.

The success of using LT-B as a carrier molecule would rely on the integral delivery of both proteins. In this study, it is observed that the ratio of fusion protein degradation is markedly different in lines P308c (P35S-BSP-LT-B::GFP) and P310 (P_{zein}-BSP-LT-B::GFP) (Fig. 2D). A higher ratio of fused protein is detected in line P310 compared to line P308c. The main difference between the two constructs is the promoter; P308c has the constitutive P35S promoter and P311 has the seed specific γ-zein promoter. As the consequence of promoter specificity, the TAEP of P308c was extracted from the stably transformed callus culture while the TAEP of P310 was from the endosperm of transgenic seed.

The presence of degraded forms of the fusion proteins is relevant from two different perspectives: (i) for localization and visualization purposes, as the truncated forms may show different localization as the uncleaved protein; and (ii) for potential implications in the use of LT-B as a carrier molecule. Because only limited numbers of callus and seed lines were analysed, it could not be determined whether this difference was due to different fusion protein processing in different tissues or due to the protein extraction process itself. More likely, however, the enhanced instability of the fusion protein is a result of stress in callus cultures, as these are not specialized for storage of proteins. Presence of a cryptic vacuolar sorting signal in the GFP sequence can result in partial degradation (daSilva et al., 2005) of fusions of mGFP5 (Haseloff et al., 1997) with a plant signal peptide. It has also been shown that certain forms of GFP can be truncated upon secretion, resulting in a lack of fluorescence (Zheng et al., 2004).

On the other hand, endosperm tissue is designed for storage and possesses all the necessary cellular machinery for correct folding, processing, and storage of proteins to high levels. Taking into account the markedly lower degree of degradation of the fusion protein in transgenic maize kernels, the results confirm that maize endosperm is an adequate tissue for recombinant protein production and accumulation.

In conclusion, this study shows that BSP, a bacterial signal peptide, can be used to direct cargo proteins to the endoplasmic reticulum of plant cells. The LT-B protein can be fused to proteins as large as 27 kDa while retaining proper folding and pentamer assembly. The fact that high fusion protein accumulation can be achieved in maize endosperm and enriched in the fibre fraction of the corn seed provide valuable information for future design strategies of specialty maize for molecular farming.

Acknowledgements

We thank François Torney, David Jackson, Martha James, Harry Horner, Gregory Phillips, and Paul Scott for helpful and stimulating discussions, Diane Bassham for a critical review of the manuscript and valuable suggestions, Margie Carter at the ISU Confocal and Image Analysis Facility and Jessica Zimmer for technical assistance, and the staff at the Maize Transformation Facility at ISU for generating transgenic material. Partial financial support for this project was provided by the United States Department of Agriculture (USDA special grant IOW05082) and the Biopharmaceutical Initiative of the Plant Sciences Institute at Iowa State University.

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