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# Expression of the 180-kD Ribosome Receptor Induces Membrane Proliferation and Increased Secretory Activity in Yeast

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**Abstract.** Expression of the canine 180-kD ribosome receptor (p180) in yeast cells resulted in a marked proliferation of intracellular membranes. The type of membranes observed varied with the expression of specific portions of p180. Rough membranes predominated when the ribosome binding domain of p180 was present, whereas expression constructs lacking this region resulted in smooth membranes. Northern analysis indicated that expression of the NH<sub>2</sub>-terminal 767 amino acids ( $\Delta$ CT), which include the ribosome binding domain, upregulated the transcription and translation of genes involved in exocytosis. The membranes that

were proliferated were functional as these cells overcame a temperature-sensitive translocation defect. Most significantly, cells that overexpressed  $\Delta$ CT and proliferated rough endoplasmic reticulum exhibited severalfold higher levels of secretion of an ectopically expressed secretory protein. We conclude that p180 expression triggers a cascade of events leading to an increase in secretory potential akin to the terminal differentiation of mammalian secretory cells and tissues.

**Key words:** secretion • endoplasmic reticulum • membrane biogenesis • yeast • ribosome receptor

**T**HE entry of proteins into the secretory pathway via the rough endoplasmic reticulum (ER) is essential for intracellular transport of proteins to the extracellular milieu as well as to a number of compartments within the cell (for review see Brodsky, 1998). Depending on their functional differentiation, different cell types possess intracellular membranes that comprise this pathway to a lesser or greater extent. Thus, in tissues with high secretory activity, such as pancreas or liver, or the plasma cells of the immune system, the secretory apparatus is developed to a level at which its membranous elements dominate the morphological landscape. A question central to both cell and developmental biology is: What is the molecular basis of the biogenesis of the secretory pathway in such tissues? This is clearly a complex problem that would benefit greatly from a simple, genetically manipulable experimental system. An interesting option has presented itself that relies on the ability of a ribosome binding protein, p180, to induce the proliferation of membranes in yeast (Wanker et al., 1995).

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Pioneering studies by Palade, Rutter, and others documented a massive proliferation of membranes during the ontogeny of hepatocytes and cells of the pancreas (Dallner et al., 1966; Wessels and Evans, 1968; Pictet et al., 1972). Through a combination of morphological and biochemical studies they showed that around the time of birth, extensive membrane proliferation is accompanied by an increase in marker enzymes. Most important is the fact that these changes coincide with the onset of the animal's ability to secrete tissue specific proteins. In B lymphocytes, similar changes in morphology and cellular physiology occur during antigen-induced maturation into plasma cells (Shohat et al., 1973). As with pancreas and liver, these changes result in a dramatic increase in secretory capacity.

The most obvious morphological change seen in liver, pancreas, and B cells is the proliferation of rough ER. Studies by Sabatini and others have shown that in such tissues the polysomes associated with the ER are predominantly synthesizing secretory products (for review see Hortsch and Meyer, 1986). These polysomes are anchored to the ER both by nascent polypeptide chains and by interactions with ER membrane proteins (Rosbash and Penman, 1971; Adelman et al., 1973; Borgese et al., 1974). One such ER membrane protein, having a molecular mass of 180 kD, was originally isolated in a functional assay from pancreatic microsomes, and has been shown to induce ri-

bosome binding both in vivo and in vitro (Savitz and Meyer, 1990, 1993; Wanker et al., 1995). It displays tissue-specific expression, with highest levels observed in typical secretory tissues, e.g., pancreas, liver, and placenta (Langley et al., 1998). This receptor, referred to here as p180, consists of three domains (see Fig. 1). At the NH<sub>2</sub> terminus is its membrane anchor. It is followed by a highly repetitive basic domain that is responsible for its ribosome binding ability (Wanker et al., 1995). In the canine as well as the human form of p180, the number of uninterrupted consensus decapeptide motifs in the ribosome binding domain is 54 (Wanker et al., 1995; Langley et al., 1998). The function of its acidic COOH-terminal domain is currently unknown, although based on the presence of typical heptad repeats, it is believed to form an  $\alpha$ -helical double-stranded coiled-coil rod (Leung et al., 1996; Langley et al., 1998). The p180 protein has a highly conserved primary structure, as the overall level of amino acid identities between the canine and human forms is 93%, rising to 99% for the NH<sub>2</sub>-terminal 86 amino acids.

A remarkable feature of p180 was first seen in studies in which the gene was expressed in yeast. When constructs encoding p180 were expressed under the control of a regulatable promoter, a rapid and significant proliferation of membranes occurred (Wanker et al., 1995). Constructs whose primary sequence included the repetitive second domain induced the proliferation of rough membranes, while those lacking this region induced smooth membrane proliferation. The expression of membrane proteins in yeast was shown previously to result in the production of membranes of varying morphologies, but never rough ER (Ohkuma et al., 1995; Koning et al., 1996). Obvious questions arising from these studies include: Are the observed membranes functional rough ER? Does the cell show an increased secretory capacity? Is what we have observed in yeast comparable to the induction of the secretory pathway in developing mammalian cells?

In this study we have ectopically expressed constructs encoding different forms of p180 in yeast. The transfected cells take on morphologies reminiscent of highly developed mammalian secretory cells. Our molecular and biochemical studies indicate that an upregulation of the secretory pathway has taken place with a striking increase in secretory capacity. These results support using yeast as a genetically manipulable model for the study of the differentiation of (mammalian) secretory cells.

## Materials and Methods

### Construction of Expression Plasmids

Regulated expression of the constructs used in this study was achieved through the use of the vector pYEX-BX that utilizes the copper-inducible promoter from the *CUP1* gene (Amrad Biotech). For the construction of the p180FL,  $\Delta$ CT, and  $\Delta$ NT we cloned BamHI-SalI fragments of the vectors pRRFL-EW1, pBSRRACT, and pBSRRANT (Wanker et al., 1995), respectively, into pYEX-BX. For the construction of pYEX-BX-AA1-151 we amplified a fragment from pRRFL-EW1 using the primers 5'-AAGGATCCATGGATATTTACGACACTCAGACC-3' and 5'-AAGTCGACTTACTCCTGGGAGCAGTTTCTAA-3'.

The fragment was cloned into pYEX-BX using BamHI and SalI sites. Plasmids were transfected into *Escherichia coli* XL1-Blue by the method of Cohen et al. (1972). The *Saccharomyces cerevisiae* strain SEY 6210 (MAT $\alpha$ leu2-3,112 ura3-52 his3- $\Delta$ 200 trp1- $\Delta$ 901 lys2-801 suc2- $\Delta$ 9) (Wils-

bach and Payne, 1993) was transfected using a lithium acetate based method (Ito et al., 1983). The copper-dependent expression of all constructs was assayed by Northern and Western blot analyses. *HMG1* and *HMG2* clones were obtained from Robin Wright (University of Washington, Seattle, WA) and subcloned into the BamHI and EcoRI sites of pYEX-BX for copper-regulated expression.

Cells were grown on SD or Sgal medium for experiments using galactose induction. SD medium, with or without 50 mM copper sulfate, was used for experiments in which proteins were expressed under control of the *CUP1* promoter. SEY 6210 was the strain used for all experiments. After 5 h of growth, transformed cells were in logarithmic phase and were used for further study.

### Electron Microscopy

Yeast cells were spheroplasted with oxalycase in SD sorbitol buffer, fixed in 2% glutaraldehyde, and postfixed with 1% OsO<sub>4</sub> in sodium cacodylate solution. Samples were dehydrated in ethanol and embedded in Spurr (Ted Pella, Inc.). Sections ~60 nm thick were made with an MT6000-XL ultramicrotome (RMC, Inc.) and stained with uranyl acetate and lead citrate. Sections were examined with a JEM-1200EX electron microscope (JEOL).

### RNA Isolation and Northern Blot Analysis

RNA isolation was performed by the method of Hollingworth et al. (1990). Total RNA (10  $\mu$ g) was separated on a 1.2% formamide containing agarose gel (Maniatis et al., 1982) and transferred to Magna Membranes (Micon Separations). Probes were generated either from restriction fragments of cloned DNA, or through PCR using the primer pairs below using yeast genomic DNA as a template: 28S RNA, 5'-TGACCTCAAATCAGGTAGGA-3' and 5'-TGTACTTGTTTCGCTATCGGT-3'; *PYK1*, 5'-AAAGATTGACCTCATTAACG-3' and 5'-GAGACTTGCAAAGTGTGGGA-3'; *KAR2*, 5'-AGTCTCAAGGGAAAAAGCGT-3' and 5'-AGCTTCCAGCAGCAAAAATT-3'; *SEC61*, 5'-ATGTCCTC-CAACCGTGTCTA-3' and 5'-AAAATCCTGGAACGAGGTTCT-3'; *OST1*, 5'-GGTATATATCATTCAAACGTG-3' and 5'-CAAGACGC-AACTACAAAA-3'; and *GDA1*, 5'-TGGCGCCCATCTTTAGA-AAT-3' and 5'-CAAGCTGATTGAATTTTAC-3'.

Northern blots were quantified using a PhosphorImager and Imagequant software (Molecular Dynamics). All lanes were corrected for loading inconsistencies by normalization to expression of *PYK1*. Values expressed are relative to vector-only controls.

### Immunofluorescence Microscopy

The antiserum to RRp has been described previously (Wanker et al., 1995). The antiserum against Sec61p was kindly provided by R. Schekman (University of California, Berkeley, Berkeley, CA) and the anti-Gda1p antisera by G. Payne (University of California, Los Angeles, Los Angeles, CA). Immunofluorescent staining of yeast cells was performed according to Adams and Pringle (1984) and Pringle et al. (1991). FITC-conjugated goat anti-rabbit antibodies were purchased from Sigma Chemical Co. The images were generated using a Zeiss fluorescence laser microscope.

### Complementation of *sec63*

The temperature-sensitive strain *ptl1/sec63* (Toyn et al., 1988) was transformed with the expression plasmids and grown at the permissive temperature on selective media. Following the induction of the expression for 4 h, 10-fold serial dilutions were plated and incubated at 24°C or 37°C for 6 or 3 d, respectively.

### Rescue from BPTI Toxicity

SEY 6210 were cotransformed with a galactose-inducible high copy plasmid expressing secreted bovine pancreatic trypsin inhibitor (BPTI)<sup>1</sup> (Parekh et al., 1995) and pYEX-BX, pYEX-BX-p180FL, - $\Delta$ NT, or - $\Delta$ CT. Following induction of copper-dependent expression of the p180 constructs, 10-fold serial dilutions were plated on glucose or galactose containing media and incubated for 3 or 5 d at 30°C. For quantification of the

1. Abbreviations used in this paper: BPTI, bovine pancreatic trypsin inhibitor; UPR, unfolded protein response.

secreted BPTI in the supernatant, aliquots were tested as described by Parekh et al. (1995).

## Results

### *p180 Expression Results in Extensive Rough Membrane Proliferation in Yeast*

Preliminary observations indicated that the expression of various domains of p180 led to membrane proliferation in yeast (Wanker et al., 1995). To refine this system for the study of membrane biogenesis, we subcloned p180-encoding constructs under control of the regulatable *CUP1* promoter (Fig. 1). High levels of expression were achieved by growing cells in copper-containing medium. Expression of the full-length ribosome receptor led to the generation of cells whose morphology (Fig. 3) was quite different from vector-only controls (Fig. 2). The cytoplasm was filled with rough (ribosome-studded) membranes that appeared in many places to have direct connections to the nuclear envelope. There was no restriction of the membranes to certain parts of the cell; rough membranes were seen in all areas between the nucleus and the periphery of the cell.

This result was quite different from the types of membranes observed in cells where proliferation had been induced by the expression of the *HMG1* gene (Wright et al.,

1988). In that case, tightly packed layers of membranes were observed in the perinuclear region, and were accordingly named karmellae. When only the first 151 amino acids of p180 were expressed, corresponding to a stretch of amino acids from the NH<sub>2</sub> terminus to the repeat region (including the membrane-spanning domain (see Fig. 1), we obtained cells with proliferated membranes identical in appearance to karmellae (Fig. 4). Many, but not all, proteins with membrane spanning domains can induce karmellae (Vergères et al., 1993; Ohkuma et al., 1995; Parish et al., 1995). The expression of soluble forms of p180, i.e., ones lacking the NH<sub>2</sub>-terminal membrane anchor (see Fig. 1), did not induce membrane proliferation (data not shown). In close agreement with previously published data (Wanker et al., 1995), the frequency with which proliferated membranes could be seen in thin sections—in all of the studies described here—was between 40 and 60%.

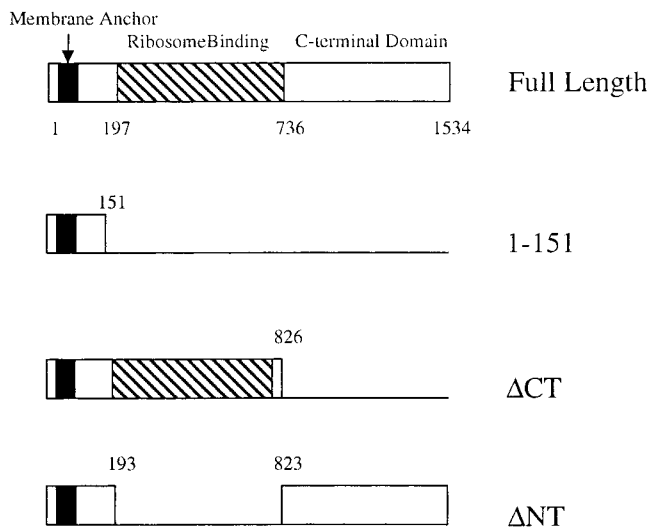
### *Expression of Different Parts of p180 Results in the Proliferation of Intracellular Membranes Having Different Morphologies*

We constructed two versions of p180 harboring deletions in major domains (Fig. 1). The first lacks the ribosome binding domain (the 54 tandem repeats of the decapeptide motif) and is referred to as  $\Delta$ NT. The second lacks the COOH-terminal coiled-coil forming domain and is referred to as  $\Delta$ CT (see Materials and Methods and Wanker et al., 1995 for details). Expression of  $\Delta$ NT resulted in the proliferation of smooth membranes, with consistent 80–100 nm spacing, from the perinuclear region to the cell periphery (Fig. 5). The smooth appearance would be expected, as the  $\Delta$ NT construct lacks the ribosome binding domain. In fact, ribosomes are selectively excluded from areas of the cell where the smooth membranes are located, and are instead restricted to more peripheral areas of the cell. This is quite different from the random distribution of ribosomes throughout the cytosol of wild-type yeast (Fig. 2). It is interesting to note, however, that the presence of the COOH-terminal domain on the  $\Delta$ NT construct results in a definitely nonkarmellar type of membrane proliferation. In this case, the spacing between membranes is far greater than in that of karmellae, and could be a feature of the ability of  $\Delta$ NT to interact through its putative coiled-coil domains with cytosolic components.

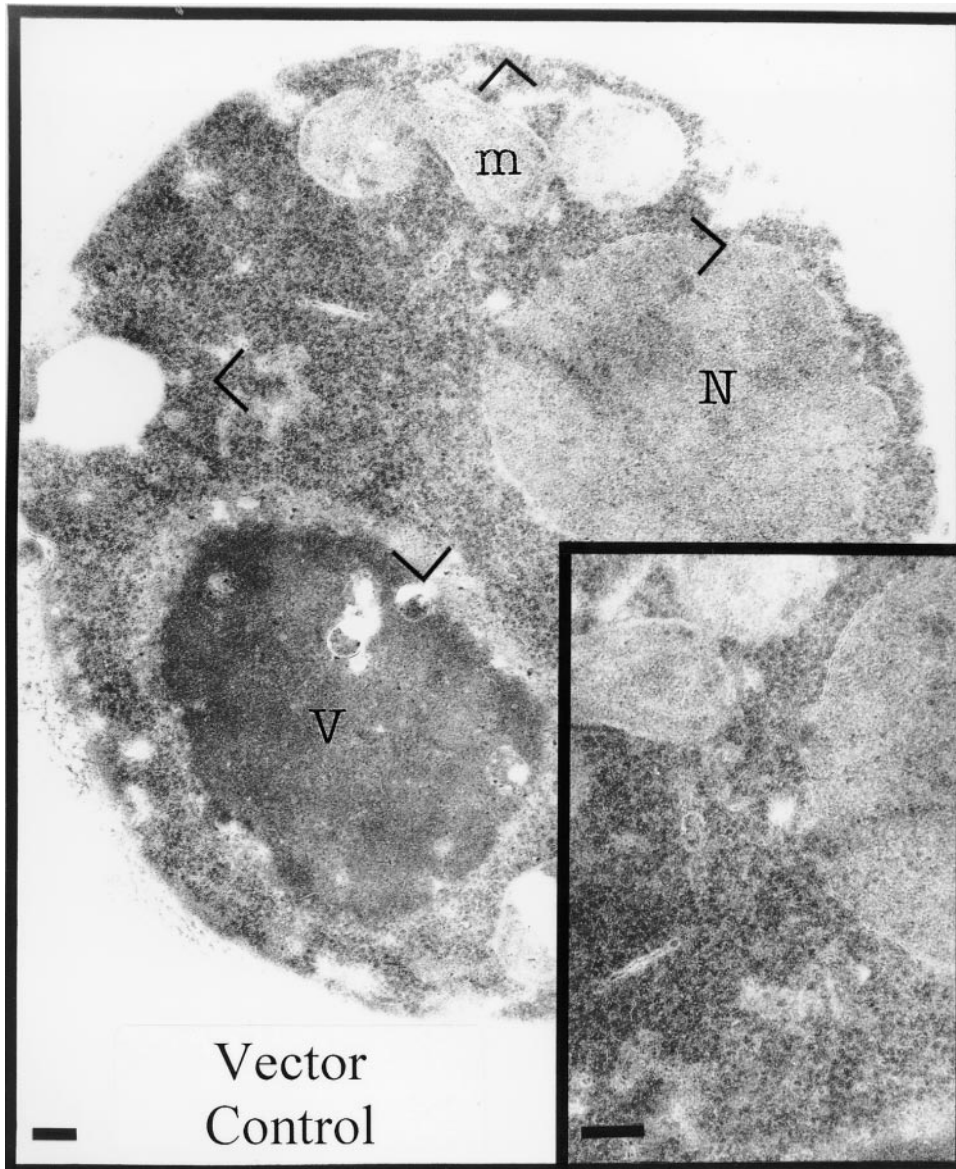
A different, yet equally striking morphology was observed in the case of the  $\Delta$ CT construct. In this case, membrane spacing collapses to that closely resembling karmellae, yet there is no restriction to the perinuclear area, and several areas of proliferated membranes are observed at the cell periphery (Fig. 6). In contrast to  $\Delta$ NT, the membranes have attached ribosomes, as indicated by the dense staining in the intermembrane space, and a lower density of free cytosolic ribosomes compared with  $\Delta$ NT.

From these data, obtained through the expression of the various domains of p180, we can make the following provisional conclusions about the morphology of the induced membranes: The presence of the NH<sub>2</sub>-terminal 151 amino acids enables membrane proliferation in general, with the closely spaced perinuclear appearance of karmellae. The presence of the repeat region enables ribosome binding irrespective of membrane distribution or spacing, and the

### p180 Constructs Used in this Study



**Figure 1.** Constructs used in this study. A description of cloning strategies can be found in Materials and Methods. The striped area, encoding a domain with 54 tandem decapeptides, has been shown previously to be essential for ribosome binding (Wanker et al., 1995). Numbers shown correspond to amino acids in the primary structure. In the text, the full-length construct is referred to as p180FL. When p1-151 is expressed, a protein is produced that comprises the NH<sub>2</sub>-terminal 151 amino acids of p180. The Membrane Anchor designation refers to amino acids 6–33, which are uncharged and mainly hydrophobic, and whose deletion leads to the production of a soluble form of p180.



**Figure 2.** Vector-only control yeast cells have few intracellular membranes. Cells were transformed with pYEX-BX. Transformants were selected and grown in  $\text{Cu}^{2+}$ -containing medium for 5 h before preparation for electron microscopy. Visible are mitochondria, the nucleus, the vacuole, and substantial quantities of free ribosomes in the cytosol. Bars, 200 nm.

presence of the COOH-terminal domain enables a large intermembrane separation.

### ***p180 Expression Results in the Coordinate Expression of Rough ER-specific Genes***

These results beg the question as to the functionality of the proliferated membranes. Are they merely lipid bilayers produced to soak up an excess of ectopically expressed membrane protein, or do they represent bona fide rough and smooth ER? The first answer to this question comes from an examination of levels of expression of genes encoding resident proteins of these membrane systems. Northern blotting (Table I) was carried out on RNA derived from p180FL,  $\Delta\text{NT}$ ,  $\Delta\text{CT}$  expressing strains, vector-only controls, and cells that overexpressed *HMG1* and *HMG2* genes. Overexpression of *HMG1* induces karmellae, whereas *HMG2* expression induces short karmellae, parallel membrane strips near the cell periphery, or membranous whorls in the cytoplasm (Koning et al., 1996). Transcripts examined included ones encoding pyruvate ki-

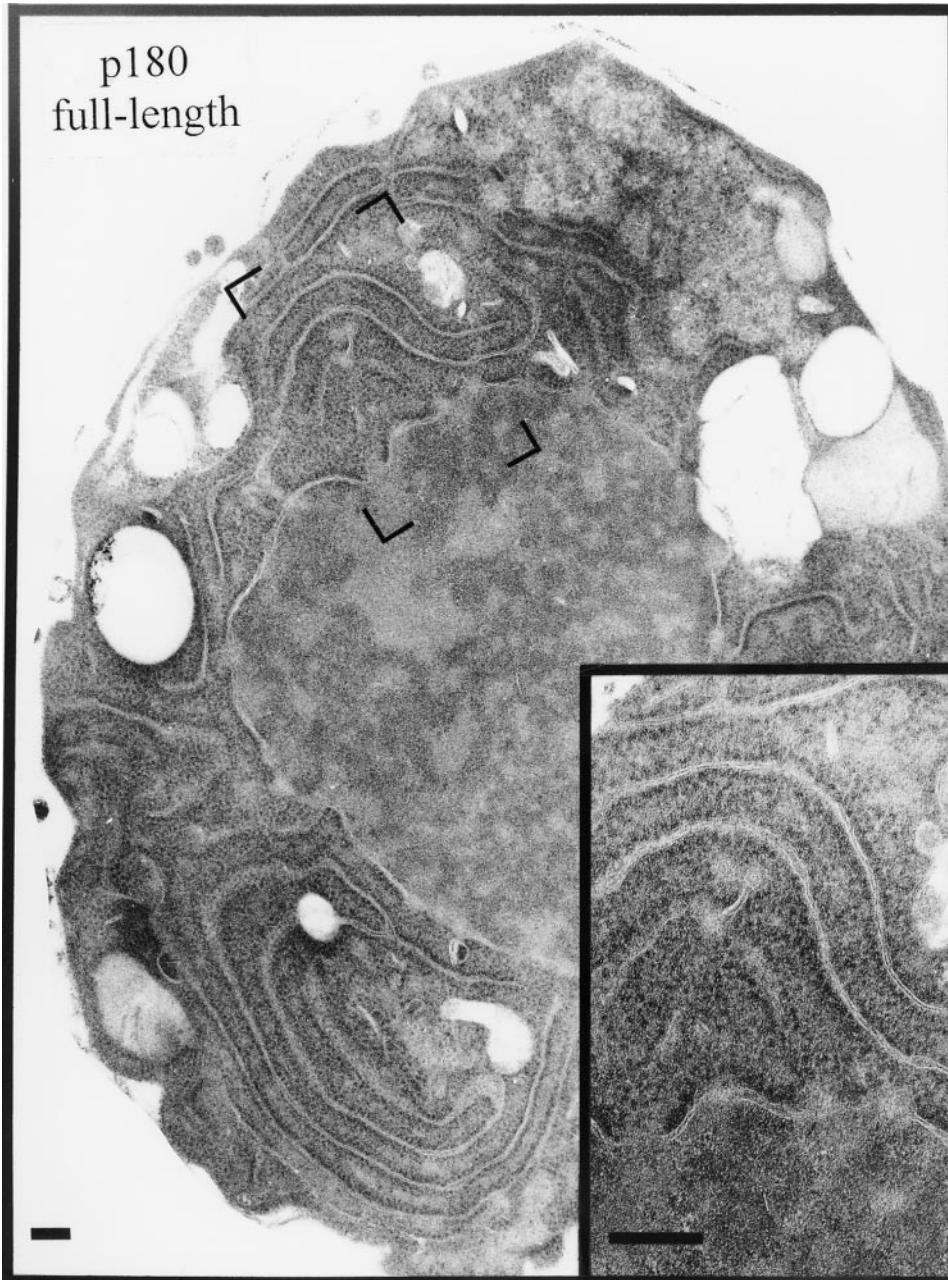
nase (*PYK1*), a cytosolic marker, and *KAR2*, *SEC61*, *SEC72*, and *OST1* that encode rough ER-specific proteins. *KAR2* encodes the luminal HSP-70 required for translocation (Vogel et al., 1990), whereas the products of

**Table I. Quantification of Northern Analysis of Gene Expression Induced by p180 Constructs**

| Expression of:          | Northern Analysis Probed with: |             |              |              |             |             |             |             |
|-------------------------|--------------------------------|-------------|--------------|--------------|-------------|-------------|-------------|-------------|
| Gene:                   | <i>PYK1</i> *                  | <i>KAR2</i> | <i>SEC61</i> | <i>SEC72</i> | <i>OST1</i> | <i>SAC1</i> | <i>GDA1</i> | <i>SEC1</i> |
| Location <sup>‡</sup> : | Cytosol                        | RER/L       | RER/M        | RER/M        | RER/M       | RER/G       | Golgi       | SV          |
| Vector-only             | 1.0                            | 1.0         | 1.0          | 1.0          | 1.0         | 1.0         | 1.0         | 1.0         |
| <i>HMG1</i>             | 1.0                            | 3.5         | 1.3          | 0.9          | 0.5         | 1.7         | 0.8         | 1.4         |
| <i>HMG2</i>             | 1.0                            | 3.7         | 1.3          | 0.5          | 0.3         | 1.1         | 0.6         | 0.8         |
| p180 Full-L             | 1.0                            | 3.2         | 10.3         | 3.5          | 5.3         | 3.9         | 2.4         | 1.7         |
| p180 $\Delta\text{NT}$  | 1.0                            | 2.2         | 2.9          | 2.7          | 4.1         | 2.7         | 1.8         | 1.2         |
| p180 $\Delta\text{CT}$  | 1.0                            | 3.8         | 14.9         | 3.6          | 6.9         | 9.1         | 5.0         | 6.1         |

\*Differences in loading were corrected using *PYK1* expression. Values shown reflect changes in expression relative to vector-only controls.

<sup>‡</sup>Abbreviations: RER/L, rough ER lumen; RER/M, rough ER membrane; RER/G, RER/Golgi; SV, secretory vesicles.



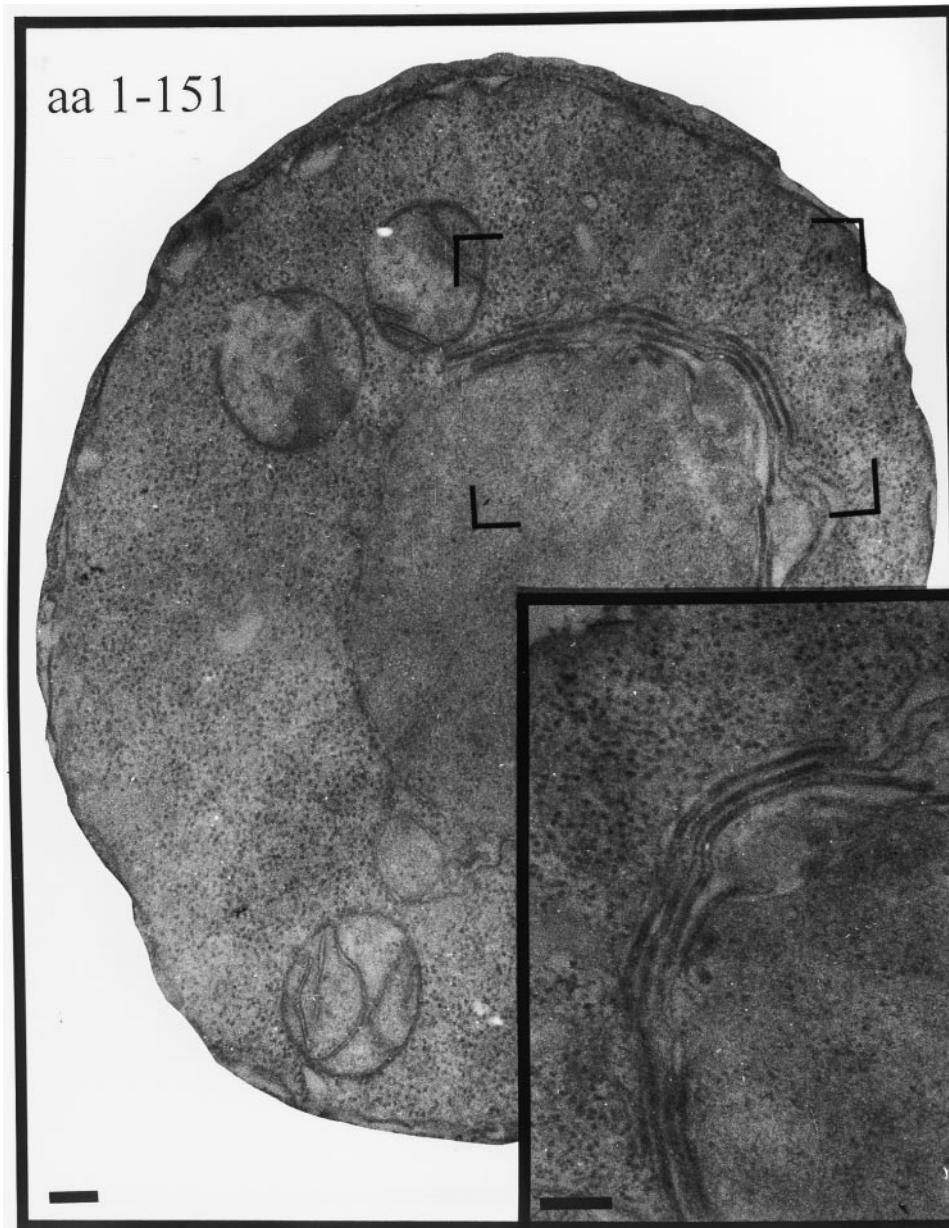
*Figure 3.* Expression of full-length p180 induces rough membrane proliferation. Cells were transformed with pYEX-BX containing full-length p180 cDNA. Transformants were selected and grown in  $\text{Cu}^{2+}$ -containing medium for 5 h before preparation for electron microscopy. Visible are extensive arrays of ribosome-studded membranes. Bars, 200 nm.

*SEC61* and *SEC72* are membrane proteins that participate in protein translocation into the rough ER (Rothblatt et al., 1994). *OST1* is part of the oligosaccharyl transferase complex, which mediates N-linked oligosaccharide addition to glycoproteins (Silberstein et al., 1995). Should any of the proliferated membranes represent real rough ER, one would predict that these markers would be induced.

Overexpression of all of the constructs resulted in an upregulation of *KAR2*, ranging from a low of 2.2-fold in the case of  $\Delta\text{NT}$  to a high of 3.8-fold for  $\Delta\text{CT}$  (Table I). In contrast, transcripts encoding rough ER membrane proteins were upregulated to the greatest extent in strains in which the ectopically expressed versions of p180 contained the ribosome binding domain (p180FL and  $\Delta\text{CT}$ ). *SEC61* was the most highly expressed of all, where close to 15-fold

higher levels were achieved in cells expressing  $\Delta\text{CT}$ . In the same strain, *SEC72* and *OST1* expression increased 3.6- and 6.9-fold, respectively. In contrast, the *HMG1 HMG2* overexpressing strains showed significant increases only in *KAR2* expression. Cells expressing the p180 construct lacking the ribosome binding domain ( $\Delta\text{NT}$ ) showed only a (comparatively) moderate upregulation of ER markers, despite high levels of membrane proliferation (see Fig. 5).

Increased levels of transcription of membrane protein genes translates into increased levels of the proteins they encode (Wanker et al., 1995). To demonstrate that these proteins are incorporated correctly into membranes, immunofluorescent techniques were used. ER-localized Sec61p was detected in the perinuclear membranes of control cells (Fig. 7 B), as is typically observed using anti-ER



*Figure 4.* Expression of amino acids 1–151 of p180 induces membrane proliferation having a karmellar morphology. Cells were transfected with pYEX-BX-AA1-151. Transformants were selected and grown in  $\text{Cu}^{2+}$ -containing medium for 5 h before preparation for electron microscopy. Visible are perinuclear membranes similar in appearance to karmellae. Bars, 200 nm.

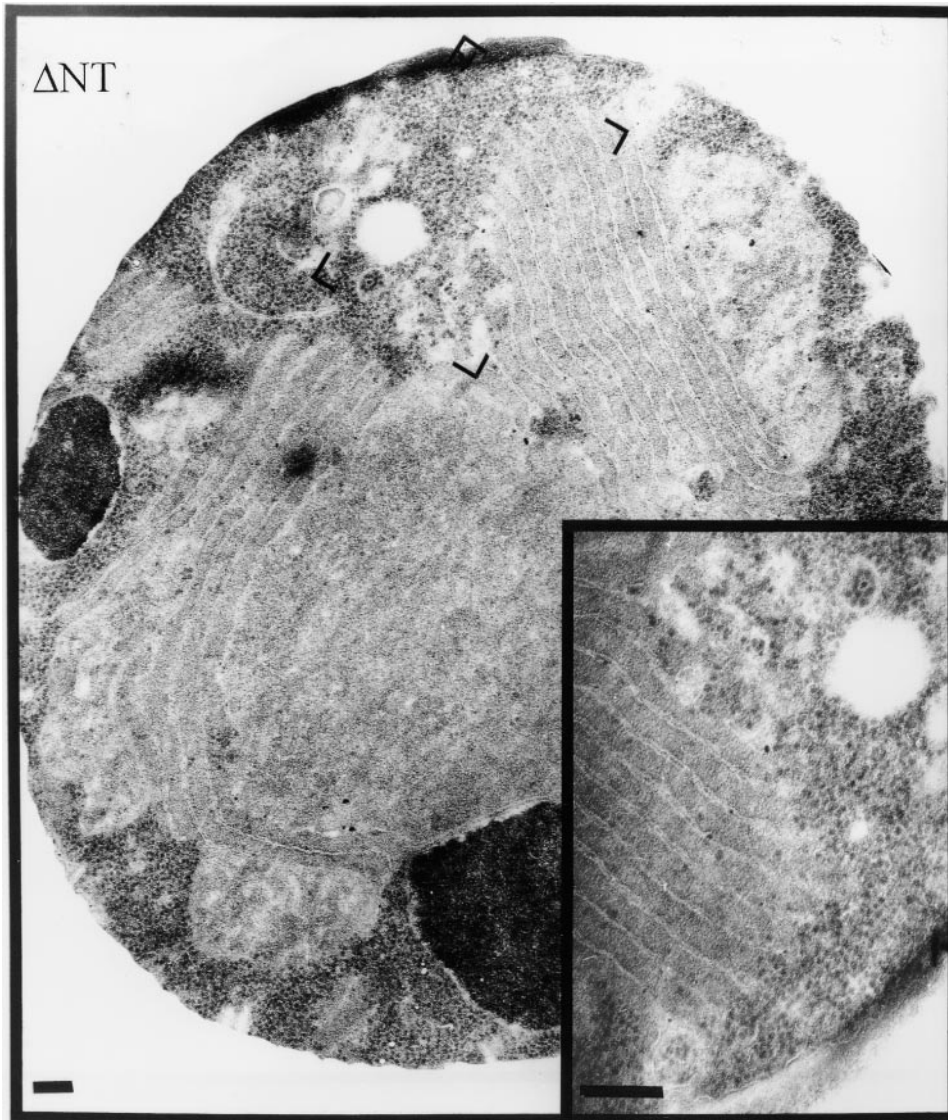
protein antibodies. Membranes proliferated in response to  $\Delta\text{CT}$  expression show a marked increase in the intensity of anti-Sec61p staining (Fig. 7 D) comparable to the detection of p180 using anti-p180 antibodies (Fig. 7 C). The localization of the fluorescence in the  $\Delta\text{CT}$  strain overlaps nicely with the location of the proliferated membranes seen in the electron microscope (Fig. 6). This result is a good predictor that functional membranes are being produced.

#### ***Rough Membranes Produced by p180 Expression Are Functional***

To establish the functionality of the proliferated rough ER, we turned to a genetic approach. We used a previously isolated strain that harbors a temperature-sensitive translocation defect. Originally isolated as *ptl1* (Toyn et al.,

1988), and subsequently shown to be capable of rescue by a wild-type *SEC63* gene (Crowe, J., and D.I. Meyer, unpublished observations), this strain has a decreased ability to translocate nascent secretory proteins *in vivo* as well as *in vitro* at the nonpermissive temperature. Although membranes produced through expression of p180 constructs would still have the defective *PTL1/SEC63* gene product, compensatory levels of translocation, and hence cell growth, should be observed due to more abundant, albeit defective, rough ER. As can be seen in Fig. 8, *ptl1/sec63* cells transfected with vector alone showed a marked reduction in growth at the nonpermissive temperature ( $37^\circ\text{C}$ ). A significant increase in cell growth at the nonpermissive temperature, compared with controls, was observed in the case of cells transfected with p180FL, and an even better rescue was observed in the case of  $\Delta\text{CT}$  expression, in which growth was virtually the same as vector-





*Figure 5.* Expression of p180 without its ribosome binding domain produces extensive smooth membrane proliferation different from karmellae. Cells were transfected with pYEX-BX containing the  $\Delta$ NT version of p180. Transformants were selected and grown in  $\text{Cu}^{2+}$ -containing medium for 5 h before preparation for electron microscopy. Visible are parallel arrays of smooth membranes. Note the exclusion of cytoplasmic ribosomes from the areas where membranes are evident, and their high density in membrane-free zones. Bars, 200 nm.

only at the permissive temperature. In contrast, membrane proliferation alone did not appear to be determinative in rescue, as expression of either *HMG1* or  $\Delta$ NT was not much better than vector-only. From these sets of experiments we conclude that the rough membranes produced in response to  $\Delta$ CT expression are functional rough ER, and, from the previous data, that membrane composition is at least qualitatively similar to wild-type.

#### ***p180 Expression Results in Coordinate Expression of Secretory Pathway Genes***

The results presented to this point indicate that functional rough ER is being produced in response to  $\Delta$ CT expression. Are the more distal aspects of the secretory pathway being induced as well? We answered this question in two ways: by Northern analysis of marker genes for organelles that function post-ER in the secretory pathway, and morphologically using immunofluorescent detection of the Golgi complex. The following markers were examined: *Sac1p*, which is involved in nucleotide transport and has

been localized to ER and to Golgi membranes; *Gda1p*, which encodes the guanosine diphosphatase required in the Golgi complex for oligosaccharide elaboration; and *Sec1p*, which is a cytosolic/peripheral membrane protein required in the exocytic fusion of secretory vesicles with the plasma membrane. The results shown in Table I demonstrate that transcripts encoding *SAC1*, *GDA1*, and *SEC1* gene products were all produced at significantly (5–10-fold) higher levels in cells expressing  $\Delta$ CT compared with controls. In contrast, inducers of smooth membranes, such as the *HMG* genes or  $\Delta$ NT expressed these markers at levels close to those of the vector control. Although one cannot test for all genes involved in secretion, these three—as well as the ER markers assessed previously—would likely participate in any process that would increase the secretory capacity of the cell.

Fluorescence microscopy on control and  $\Delta$ CT expressing cells was performed to further characterize the expression of post-ER markers. We used an anti-*Gda1p* antibody to reveal the Golgi complex in these cells. As can be seen in Fig. 9 A, a single Golgi body can be visualized in vector-





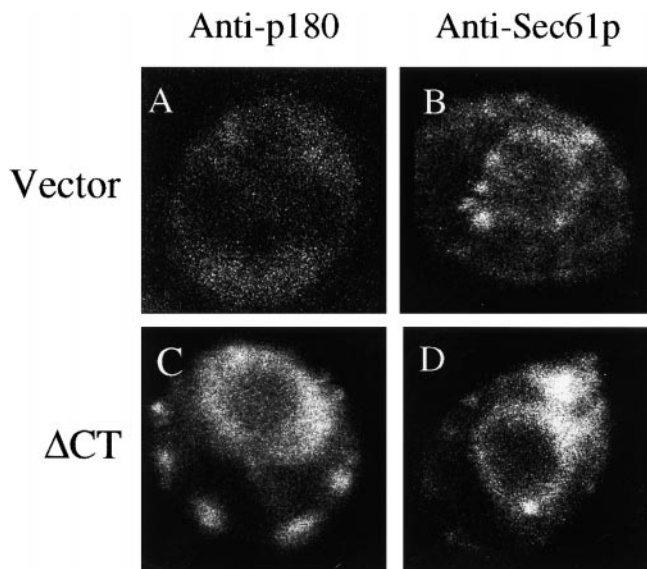
*Figure 6.* Expression of the NH<sub>2</sub>-terminal half of p180 induces extensive, closely spaced rough membranes. Cells were transfected with pYEX-BX containing the ΔCT version of p180. Transformants were selected and grown in Cu<sup>2+</sup>-containing medium for 5 h before preparation for electron microscopy. Visible are closely opposed membranes with a lower density of ribosomes in membrane-free zones (compared with cells shown in Fig. 4). Bars, 200 nm.

only cells. In contrast, numerous Golgi appear in ΔCT-expressing cells (Fig. 9 B), some larger and some smaller. Taken together, these data suggest that the entire secretory pathway may be upregulated in response to ΔCT expression.

#### ***p180 Expression Results in Increased Secretory Capacity***

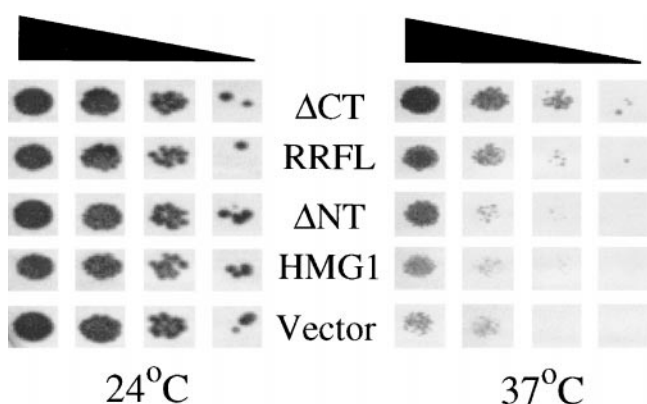
To get an approximation of the secretory capacity of wild-type yeast, we transfected control strains with a plasmid encoding BPTI. BPTI is a low molecular weight protein that can be easily measured colorimetrically when it ap-

pears in the growth medium. The assay is based on the ability of BPTI to inhibit trypsin hydrolysis of an artificial substrate. Accordingly, BPTI was expressed under *GAL* control in glucose or in galactose-containing medium. Interestingly, once levels of BPTI secretion approach 8–10 μg/ml in wild-type cells grown under standard culture conditions, cell growth is inhibited, ostensibly by blocking or overwhelming the secretory process (Parekh et al., 1995). Fig. 10 shows that vector-only or the smooth membrane-proliferating ΔNT cells were unable to grow when BPTI was expressed through galactose induction for a period of 24 h. On the other hand, cells expressing BPTI were rescued by the expression of p180FL or ΔCT. Our prelimi-

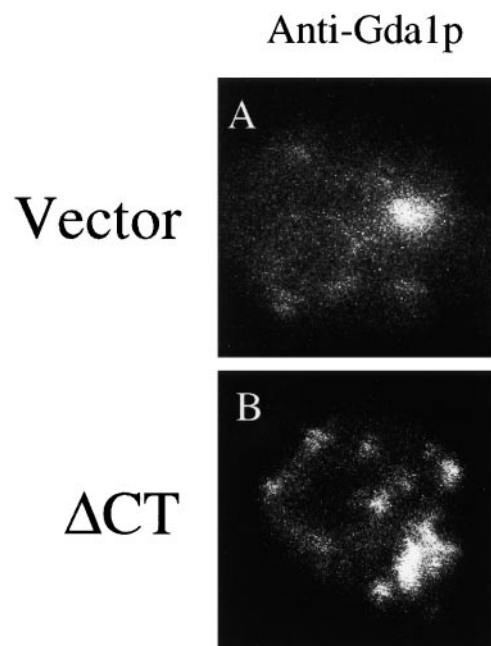


**Figure 7.** Expression of  $\Delta$ CT results in a proliferation of Sec61p-containing membranes. Cells were transfected with pYEX-BX containing the  $\Delta$ CT version of p180 cDNA. Transformants were selected and grown in  $\text{Cu}^{2+}$ -containing medium for 5 h before preparation for immunofluorescence microscopy. (A and C) Vector-only and  $\Delta$ CT-expressing cells stained with anti-p180 antibodies. (B and D) Vector-only and  $\Delta$ CT-expressing cells stained with anti-Sec61p antibodies.

nary conclusion was that an increase in secretory capacity enabled toxic quantities of BPTI to be removed from the cells. Recent studies by Wittrup and co-workers (Shusta et al., 1998) identified luminal proteins (Kar2p and PDI) whose increased expression enabled higher levels of single chain antibody secretion. These findings are consistent with the results presented here, as the expression of p180FL or  $\Delta$ CT induces increases in levels of transcripts



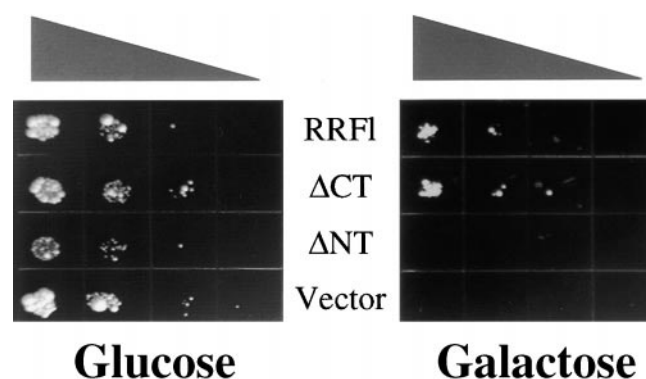
**Figure 8.** Induced membrane proliferation rescues a temperature sensitive translocation defect. Cells harboring the *sec63/ptl1 $\Delta$*  mutation were transformed with the constructs shown in the middle of the figure (RRFL is the same as p180FL). Transformants were serially diluted (10-fold steps) onto  $\text{Cu}^{2+}$ -containing medium and grown at either the permissive (24°C) or nonpermissive (37°C) temperature. The triangle above the figure indicates decreasing cell concentrations that were plated.



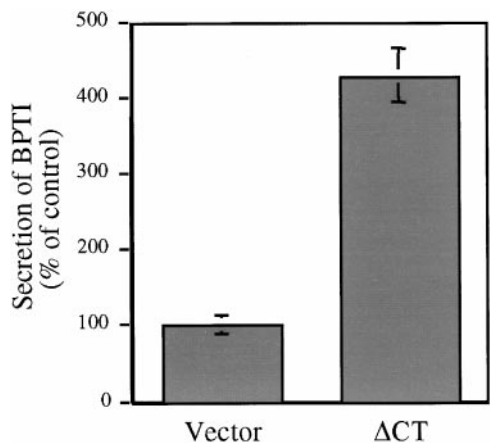
**Figure 9.** Expression of  $\Delta$ CT results in a proliferation of Gda1p-containing membranes. Cells were grown and prepared as described in Fig. 7. Cells were stained with antibodies against the Golgi marker protein Gda1p.

encoding these proteins (Table I and Becker, F., unpublished results).

Measuring BPTI accumulation in the medium substantiated the hypothesis that the toxic effect of BPTI was overcome through increased secretion. In the case of cells grown on glucose, BPTI secretion was undetectable. In the case of cells expressing  $\Delta$ CT and grown on galactose, levels of BPTI that accumulated in the medium during the assay rose >400% compared with control cells (Fig. 11). In contrast, membrane proliferation alone—as observed with



**Figure 10.** Expression of full-length p180 or  $\Delta$ CT rescues BPTI-induced growth arrest. Cells transformed with the constructs indicated in the middle of the figure (RRFL is the same as p180FL) were subsequently transformed with a plasmid directing the expression of BPTI under galactose control, and serially diluted (10-fold steps) onto  $\text{Cu}^{2+}$ -containing growth medium that included either glucose or galactose as the carbon source. Triangle above figure indicates decreasing cell concentrations that were plated.



**Figure 11.** Expression of  $\Delta$ CT significantly increases BPTI secretion. Control and  $\Delta$ CT-expressing cells, prepared as described in the legend to Fig. 10, were grown on galactose and copper-containing medium. BPTI secreted into the medium was determined by the colorimetric assay described in Materials and Methods.

$\Delta$ NT expression—affected neither rescue from BPTI-mediated growth arrest (Fig. 10), nor BPTI levels in the medium (data not shown). We take this to be direct proof of an increased secretory capacity mediated through the expression of  $\Delta$ CT.

## Discussion

Our findings of an explosive proliferation of functional membranes coupled to an overall increase in secretory capacity in response to the heterologous expression of a single protein raise a number of intriguing questions about the regulation of membrane biogenesis. Previous reports, spanning more than three decades, have documented smooth membrane proliferation as a physiological response to the administration of xenobiotics (Orrenius et al., 1965; McLaughlin et al., 1999) or resulting from the overexpression of specific ER membrane proteins in mammalian cells (Anderson et al., 1983). To a certain extent, these processes have been characterized in molecular detail. In yeast, the overexpression of *HMG1* and *HMG2* genes in *S. cerevisiae* produced a number of smooth, closely spaced membranes whose specific morphologies differed depending on the isoform that was overexpressed. In these cases membranes appeared either as karmellae, short karmellae, strips, and/or whorls (Koning et al., 1996). In *C. maltosa*, the overproduction of cytochrome P450 produced smooth ER tubules as well as karmellae (Ohkuma et al., 1995). Sec61p, a component of the translocation pore, appeared to be a component of the proliferated membranes (Wiedmann et al., 1993; Koning et al., 1996) indicating their derivation from the ER. However, neither an apparent accumulation of rough membranes, nor an increase in secretory activity was observed.

In the case of p180, a strikingly different morphology was observed, depending upon which part of p180 was expressed. Expression of the full-length receptor resulted in a proliferation of widely spaced rough membranes. The 80–100 nm spacing correlated with the presence of the

COOH-terminal half of p180, as the  $\Delta$ NT construct induced smooth membranes with a considerable distance between them. In contrast, expression of  $\Delta$ CT as well as p1-151 produced rough and smooth membranes, respectively, both possessing a very tightly packed appearance. Based on these data, one could make the prediction that the three separate domains induce three different aspects of rough membrane biogenesis. Production of membranes per se requires the expression of a membrane anchor and some minimum number of cytoplasmically exposed amino acids, e.g., p1-151. Adding on a ribosome binding domain, as in  $\Delta$ CT, triggers the induction of functional rough ER, while addition of the COOH-terminal domain, as in p180FL and  $\Delta$ NT, results in the typical uniformly spaced parallel arrays of membranes seen in mammalian secretory tissues (Fawcett, 1981).

Parrish et al. (1995) determined that the sequence of Hmg1p that induces karmellae resides in a lumenally disposed loop located between two transmembrane helices. It is interesting to note that the primary structure of p180 predicts that fewer than half a dozen amino acids could be located in the lumen of the ER, making a comparable mechanism unlikely. This is similar to the topology of another membrane-inducing protein, cytochrome P450. In this case, there are maximally 15 amino acids that are likely exposed to the lumen (Menzel et al., 1997).

The observation that p180 seems to reach its highest expression levels in tissues with high secretory capacity (Langley et al., 1998) provides a key insight into its probable physiological role: sequestration of ribosomes to the region of the cell where substantial amounts of synthesis and transport are occurring. This hypothesis is consistent with a number of relevant findings. First shown a number of decades ago, and repeatedly observed, is the fact that roughly half of all ribosomes bound to microsomal membranes isolated from secretory tissues can be released by high salt in the absence of the nascent chain release by puromycin (Adelman et al., 1973). The implication is that such ribosomes are not involved in the synthesis of secretory or membrane proteins. p180 is a likely candidate for mediating the attachment of such uninvolved ribosomes to the ER membrane. Additionally, depletion of p180 from reconstituted translocationally competent pancreatic microsomes has a significant effect on protein translocation (Savitz and Meyer, 1993), yet it is not required for translocation when only purified components are incorporated into liposomes (Görlich and Rapoport, 1993). The high level and efficiency of in vitro protein translocation seen in intact microsomes would be due to the assistance in ribosome binding provided by p180. On the other hand, stabilization of the ribosome-membrane interaction may be transient or nonessential (Nicchitta and Zheng, 1997), or may be mediated by the Sec61-containing translocon (Kalies et al., 1994; Powers and Walter, 1998) in the highly reconstituted systems.

Why is there no p180 in yeast? The answer to this comes from the same data alluded to above. Yeast cells do not have the secretory capacity of mammalian pancreas or liver. Therefore, just as many mammalian cell types exist with minimal or no expression of p180 (Langley et al., 1998), so can yeast. Ribosome binding to the ER membrane would be achieved through the nascent polypeptide

and by interactions between the ribosome and components of the Sec61 complex (Powers and Walter, 1998).

Equally if not more intriguing is the question of how the expression of p180 results in the proliferation of membranes in the secretory pathway in a cell in which it is not normally expressed. Hypotheses can be based on the elements of the protein mentioned previously, and their functional properties. It is clear that the membrane anchoring domain plus a few dozen amino acids of the cytosolic domain induces lipid bilayer proliferation. One can expect that the synthetic machinery needed for lipid biosynthesis is switched on in these cases via transcription of the requisite enzymes. This postulate is borne out by studies on other membrane proteins (Menzel et al., 1997). Moreover, our own preliminary results show increased *INO1* expression in all of our strains in which membrane proliferation occurred (Block-Alper, L., unpublished observations). However, expression of the membrane anchor domain (p1-151) or  $\Delta$ NT did not induce *SEC61* or any of the other ER resident proteins; their expression correlated with expression of the ribosome binding domain. How then does the addition of this domain make such a difference in the number of genes that are being upregulated, and is the process dependent upon its ability to bind ribosomes with high affinity? It is tempting to speculate that the cell erroneously mobilizes greater secretory capacity due to a loss of free ribosomes from the cytosol, and that sensing this loss is the key step in induction of the relevant genes. If this is true, one could expect that the high level of expression of any secretory or membrane protein would induce the upregulation of secretion as long as it removes ribosomes from the cytosolic pool and directs them to the membrane. This has not yet been systematically investigated. On the other hand, Hanein et al. (1996) have demonstrated that Sec61p complexes form oligomeric rings in the membrane, and that their formation is dependent upon ribosome binding. A recruitment of ribosomes to the membrane could result in the formation of Sec61p complex oligomers and the corresponding decrease in free Sec61p complexes would prompt the upregulation of genes inducing membrane proliferation. These hypotheses form the basis of future experimentation.

Recent studies have elucidated an elaborate signaling pathway in rough ER. In response to a buildup of unfolded or misfolded proteins, a cascade is triggered that results in increased chaperone production (for review see Sidrauski et al., 1998). This unfolded protein response (UPR) is also linked to membrane biogenetic events, including membrane lipid production (Cox et al., 1997). The overexpression in UPR-deficient cells of certain membrane proteins, such as Hmg1p, has been shown to be lethal (Cox et al., 1997). On the other hand, UPR-deficient cells that overexpress cytochrome P450 are viable (Menzel et al., 1997). The p180-induced response also appears independently of the UPR as demonstrated by the fact that both viability and p180-induced membrane proliferation were observed in strains harboring a deletion of the *IRE1* gene (Becker, F., unpublished observations).

The work described here makes use of the canine p180 cDNA that was cloned in our laboratory. Extensive data are now available on the human p180 homologue (Basson et al., 1996; Langley et al., 1998), as well as a potential al-

ternatively spliced transcript derived from the p180 gene that was originally characterized in chicken (Krug et al., 1995), but also occurs in humans (Langley et al., 1998). Some of the transcripts processed in this way lack the repeat region entirely, yet retain the membrane anchor and the COOH-terminal region. The overall levels of similarity between canine and human p180s are >90%. The function of the splice variants is still largely speculative. Of interest is the fact that the consensus sequence of the decapeptide repeats is highly conserved, as well the total number, 54 (Wanker et al., 1995; Langley et al., 1998). The report on the human gene also points out that the COOH-terminal domain, for which function has yet to be determined—but from these studies is implied to generate the regular intermembrane spacing observed in secretory tissues—has the hallmarks of forming a coiled-coil structure. Should spacing depend upon specific interactions with elements that organize the cytoplasm, this type of secondary structure of the COOH terminus may be advantageous. On the other hand, the COOH-terminal domain could function in the regulation of ribosome binding to the repeats, or in the spatial arrangement of the molecule on the cytoplasmic surface of the rough ER. It is interesting to note that GST fusions containing the COOH terminus of p180 can bind to columns containing nucleoside triphosphates such as ATP (Savitz, A., and D. Meyer, unpublished observations), consistent with previous observations that p180 is an ATP binding protein (Zimmerman and Walter, 1991).

Elucidation of the mechanism of membrane induction by p180 will benefit greatly from the observations made here in the yeast system. Through screening strategies it should be possible to identify genes whose products are capable of stimulating the upregulation of genes essential for membrane biogenesis. Moreover, similar schemes will enable mutant strains to be identified that lack the ability to upregulate membrane biogenesis in response to p180 expression. Identification and characterization of such genes and their products should provide the required threshold for further analysis of this interesting eukaryotic regulatory pathway.

It will also be interesting to analyze rough ER membrane induction by p180 in mammalian cells. Preliminary studies show that increasing cellular levels of p180 significantly stimulates rough membrane induction in nonsecretory cells (Castro-Vargas, E., F. Becker, and D.I. Meyer, unpublished observations). This implies that the expression of p180 may play a central role in the terminal differentiation of secretory tissues. The fact that a similar phenomenon is observed in mammalian cells makes the use of the genetically manipulable yeast model described here especially attractive.

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