Oxysterol binding protein (OSBP) is the only protein known to bind specifically to the group of oxysterols with potent effects on cholesterol homeostasis. Although the function of OSBP is currently unknown, an important role is implicated by the existence of multiple homologues in all eukaryotes so far examined. OSBP and a subset of homologues contain pleckstrin homology (PH) domains. Such domains are responsible for the targeting of a wide range of proteins to the plasma membrane. In contrast, OSBP is a peripheral protein of Golgi membranes, and its PH domain targets to the trans-Golgi network of mammalian cells. In this article, we have characterized Osh1p, Osh2p, and Osh3p, the three homologues of OSBP in *Saccharomyces cerevisiae* that contain PH domains. Examination of a green fluorescent protein (GFP) fusion to Osh1p revealed a striking dual localization with the protein present on both the late Golgi, and in the recently described nucleus-vacuole (NV) junction. Deletion mapping revealed that the PH domain of Osh1p specified targeting to the late Golgi, and an ankyrin repeat domain targeting to the NV junction, the first such targeting domain identified for this structure. GFP fusions to Osh2p and Osh3p showed intracellular distributions distinct from that of Osh1p, and their PH domains appear to contribute to their differing localizations.

INTRODUCTION

The mechanisms underlying the synthesis and uptake of sterols by eukaryotic cells are now relatively well characterized. However, much less is understood about how cells regulate their intracellular sterol levels, and how they maintain a nonhomogenous distribution of sterols between different internal membranes. Sterol homeostasis requires that there must be mechanisms to sense cellular sterol levels, and although there has been much recent progress in identifying some of the key regulators of cholesterol metabolism (Brown and Goldstein, 1999), less is understood about how sterol sensing occurs. The intracellular traffic of cholesterol appears to be important in this feedback (Lange and Steck, 1996). The majority of cholesterol is found in the plasma membrane, but it is in the endoplasmic reticulum (ER), which itself has low levels of cholesterol, where the changes in cellular cholesterol levels are responded to by the sterol regulatory element-binding protein (SREBP) system that controls the transcription of genes encoding cholesterol biosynthetic enzymes (Brown and Goldstein, 1999; Lange et al., 1999). Although it might be expected that the systems controlling cholesterol metabolism would recognize cholesterol itself, there has been a long-standing interest in the possibility that oxysterols, a group of oxidized derivatives of sterols, are important second messengers in sterol homeostasis (Brown and Goldstein, 1974; Kandutsch and Chen, 1974; Accad and Farese, 1998). Indeed, oxysterols such as 25-hydroxycholesterol are up to a 1000 times more potent than cholesterol itself as down-regulators of cholesterol synthesis (Kandutsch et al., 1978; Goldstein and Brown, 1990).

Because many oxysterols can be generated readily from the nonenzymatic oxidation of cholesterol, their physiological relevance has until recently been uncertain (Smith, 1996). However, there is now increasing evidence that oxysterols play important roles in vivo. First, intracellular hydroxylases have been discovered that can convert cholesterol into specific oxysterols, including 25-hydroxycholesterol (Lund et al., 1998, 1999; Russell, 2000). Second, it has recently been shown that the enzymes responsible for the hepatic conversion of excess cholesterol into bile acids are regulated by a nuclear hormone receptor (LXRα) that binds a specific subset of oxysterols, in particular 24-hydroxycholesterol, which are synthesized when cholesterol levels rise (Janowski et al., 1996). Third, although most mammalian cells export cholesterol to high-density lipoprotein particles in the plasma, at least two cell types, macrophages and neurons, export the bulk of sterol as 27- and 24-hydroxycholesterol, respectively (Bjorkhem et al., 1999).
If intracellular oxysterols serve as second messengers there must be particular proteins that recognize them. The only protein known to bind specifically to the group of oxysterols that are active in the down-regulation of cholesterol synthesis is oxysterol binding protein (OSBP) (Dawson et al., 1989). OSBP was identified as being the most abundant cytosolic protein that bound to such regulatory oxysterols (Taylor et al., 1984; Dawson et al., 1989b). Characterization of mammalian OSBP showed that it is associated with the periphery of the Golgi and other intracellular membranes, and that this Golgi localization is stimulated by the presence of oxysterols (Ridgway et al., 1992). The function of OSBP is unclear, but overexpression in tissue culture cells has multiple effects on cholesterol homeostasis and sphingolipid synthesis (Lagace et al., 1997, 1999).

Although the precise function of OSBP has remained elusive, it at least seems certain that this function is required in all eukaryotes, because multiple OSBP homologues have been found in the genomes of all eukaryotes so far examined. These proteins all share a conserved 400 amino acid domain found at the C terminus of OSBP, which has been shown to bind oxysterols (Ridgway et al., 1992). For convenience we will refer to this shared, characteristic, domain as the “oxysterol binding domain,” although its binding specificity in other species has not been investigated. The existence of multiple OSBP homologues raises the question of whether the different proteins in a given organism have related but distinct functions. Some evidence that this is the case comes from the fact that OSBP homologues can be divided into two general classes, ones that comprise an oxysterol binding domain alone, and longer ones such as OSBP itself that have a pleckstrin homology (PH) domain at the N terminus. Most PH domains in other proteins direct localization to the plasma membrane, often by interaction with phosphatidylinositol phosphates (PIPs). We have found that, in contrast, the PH domain of OSBP specifies targeting to the trans-Golgi network (TGN) of mammalian cells, and this interaction requires the presence of Golgi PIPs (Levine and Munro, 1998).

To learn more about the functional relevance of the intracellular targeting of oxysterol binding proteins, we have studied the situation in the yeast Saccharomyces cerevisiae, which contains seven OSBP homologues (OSH genes) (Beh et al., 2001). Three of these genes (OSH1, OSH2, and OSH3) encode proteins that, like OSBP, have a large N-terminal region that includes a PH domain (Figure 1). Of these, Osh1p and Osh2p also have three ankyrin repeats, which are not found in the mammalian protein. We were interested in whether the presence of this N-terminal extension reflected a common site of action in the cell. We report here that green fluorescent protein (GFP) fusions to the three proteins are located to different parts of the cell. In particular Osh1p has a striking dual localization being found on both the Golgi and the nucleus-vacuole (NV) junction, a recently described specialization of these two organelles (Pan et al., 2000). Targeting of Osh1p to the Golgi depends upon the PH domain, whereas targeting to the NV junction is specified by the ankyrin repeat region, the first targeting domain identified for this unusual structure. This suggests that OSBP homologue function is required in multiple parts of the cell, and that as a consequence the different members of the family contain distinct targeting determinants.

![Figure 1. Structure of OSBP and its homologues in S. cerevisiae. The PH domain (PH), ankyrin repeats (Ank), and the portions of the membrane containing the oxysterol binding domain alone, are marked.](image)

### Human OSBP

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### Yeast OSBP Homologues

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**Figure 1.** Structure of OSBP and its homologues in *S. cerevisiae*. The PH domain (PH), ankyrin repeats (Ank), and the portions of the membrane containing the oxysterol binding domain alone, are marked.

### MATERIALS AND METHODS

#### Strains and Plasmids

The genotypes of yeast strains used are listed in Table 1. Deletions of OSH1, OSH2, and OSH3 were made by homologous recombination with the use of promoter and terminator fragments, between which were placed dual loxP sites flanking HIS3, LEU2, and URA3 for OSH1, 2, and 3, respectively (Sauer, 1994). In some cases markers were excised from strains containing OSH gene deletions by expression of Cre recombinase from the GAL1/10 promoter (CEN, TRP1 plasmid, growth for 3 h with galactose), followed by plating on nonselective plates and then replica plating to identify colonies in which markers were lost. Loss of markers without unwanted chromosomal rearrangement around the loxP sites was confirmed by polymerase chain reaction (PCR).

NVT1 and VAC8 were deleted with the use of the PCR method with the heterologous marker gene Schizosaccharomyces pombe HIS5 (Wach et al., 1997). GFP-Osh2p was constructed in a Δosh1 background (strains TLY227 and TLY228) by a similar integration of a PCR product comprising Kluyveromyces lactis URA3 (Langle-Rouault and Jacobs, 1995). PHOS5 promoter, GFP and a Myc epitope tag upstream of the first codon of the OSH2 open reading frame (ORF), generated with the use of pTLUPG as a template.

Plasmids used are listed in Table 1. OSH1-3 open reading frames were cloned by allele rescue with the use of flanking promoters and terminators (Rothstein, 1991), and checked by restriction digest mapping and sequencing of junctions. PHspecter was sequence as used previously (Levine and Munro, 1998). Linkers that resulted from the restriction sites between the individual components of the chimeric proteins described in Table 1 were as follows (in single-letter amino acid notation, with residues numbered according to their position in the relevant ORF): GFP to Osh1p (pTL311) = LPG; GFP to Osh2p (pTL312) = LGAGA; GFP to Osh3p (pTL313) = SNSLG; Osh1p to GFP and Osh3p to GFP (pTL321, pTL322, pTL323, pTL327, pTL335, pTL338, and pTL362) = TKLP-VTSVPEK; flanking PHspecter and PHspecter (pTL322, pTL323, pTL325, pTL331, and pTL356) = KLGS at N terminus, KNS at C terminus; Osh1p to GFP (in pTL324) = HKLGSTKLP-VTSVPEK; GFP to Osh1p to GFP (pTL325) = LGS; GFP to Osh2p to GFP (pTL342 and pTL343) = L; Osh2p to GFP (in pTL342,
RESULTS

The Three Full-Length OSBP Homologues in Yeast Are Functionally Distinct

Osh1p was previously identified as a full-length yeast homologue of mammalian OSBP (although it was initially...
Osh Proteins Have Distinct Intracellular Distributions

We next investigated the intracellular localization of Osh1p, Osh2p, and Osh3p by tagging the proteins at their N termini with GFP. When the GFP-Osh1p construct was expressed in JRY6326, a strain in which all seven OSH open reading frames are deleted but which is rescued from lethality by a copy of Osh2p expressed under the MET3 promoter (Beh et al., 2001), was transformed with an empty plasmid (pRS416); or with plasmids expressing either untagged Osh1p (pTL301), Osh1p tagged at the N terminus with GFP (pTL311); or GFP-tagged constructs containing portions of Osh1p expressed from the promoter stated in brackets. Osh1OBBD (1–690) was tagged at the C terminus with GFP and was expressed from either the OSH1 promoter (pTL362) or the TPI1 promoter (pTL321); OBD1 (689–1190) was tagged at the N terminus with GFP, and expressed from either the OSH1 promoter (pTL371) or the PHO5 promoter (pTL376). By quantitative protein blotting with anti-GFP the expression levels of the four constructs were, respectively, 0.3-, 20-, 0.7-, and 17-fold that of GFP-Osh1p. Growth conditions were as described in MATERIAL AND METHODS. Note that in the absence of OSH1, very slight growth was occasionally seen on low tryptophan.

(C) In vivo activity of GFP fusions to Osh2p and Osh3p. JRY6326, a strain in which all seven OSH open reading frames are deleted but which is rescued from lethality by a copy of Osh2p expressed under the MET3 promoter (Beh et al., 2001), was transformed with an empty plasmid (pRS416); or with plasmids expressing either untagged Osh2p and Osh3p either untagged or GFP-tagged (pTL302, pTL303, pTL312, pTL313). Growth of JRY6326 was inhibited by repression of the MET3 promoter, but GFP tagged Osh2p and Osh3p were able to rescue the lethality as effectively as the untagged proteins.
a strain in which the only other OSH gene was under a regulated promoter, the fusions were able to maintain growth when this promoter was repressed, indicating that these GFP fusions retain functional activity (Figure 2C).

When live yeast expressing the GFP fusions were examined by fluorescence microscopy in midlog phase, GFP-Osh1p was localized in punctate structures that are discussed in detail below (Figure 3A). In contrast, GFP-Osh2p was apparently localized to the plasma membrane, concentrated in the budding area of G1 phase cells, and around the mother-daughter bud-neck of S phase cells, as well as in a diffuse cytoplasmic pool (Figure 3B). At higher levels of expression, GFP-Osh2p remained localized to the plasma membrane, although it then redistributed around the whole plasma membrane (our unpublished results). GFP-Osh3p was apparently diffusely distributed throughout the cytoplasm (Figure 3C), although it is possible that there may be a functional targeted population masked by this cytosolic pool. These results show that the three full-length homologues of OSBP in yeast are spatially distinct, which suggests that they may have differing roles in the cell.

Osh1p Localizes to Two Separate Organelles

Most of the punctate structures labeled by GFP-Osh1p were small and randomly distributed throughout the cytoplasm, a pattern typical of yeast Golgi membranes (as confirmed below). However, an unusual single linear structure could also be seen at one side of the vacuole (arrows in Figure 3A). The linear structures appeared in multiple adjacent confocal sections, hence they are disk-shaped, and were occasionally seen as discs lying above or below the vacuole in the horizontal plane (our unpublished results). When live cells expressing GFP-Osh1p were labeled with DAPI to detect DNA, and with FM4-64 to visualize the vacuole, these linear structures were found to be located directly between the vacuole and the nucleus (Figure 4A). Although most cells had a single linear structure, some cells with multilobed vacuoles contained two linear structures between the nucleus and two separate vacuoles, but not between the vacuoles themselves. This distribution is the same as the recently described NV junction, a region of close contact between the nuclear envelope and the vacuole whose formation requires the ER membrane protein Nvj1p and the vacuole protein Vac8p (Pan et al., 2000).

To confirm that the linear structures containing GFP-Osh1p are NV junctions, the gene NVJ1 was deleted, because loss of this gene has been shown to lead to the loss of the NV junction (Pan et al., 2000). In Δnvj1 cells, which still contain both nucleus and vacuole, GFP-Osh1p was no longer found between these two organelles, although it was still located in punctate structures (Figure 4B). These observations were strengthened by our finding of conditions under which the only localization of Osh1p is to the NV junction. When cells were examined at stationary phase we observed a complete loss of localization to the multiple punctate structures, but a clear preservation of staining of the NV junction (Figure 4C). The basis of this altered distribution is unknown, but it does not appear to reflect dispersal of the Golgi because a GFP fusion to the late-Golgi t-SNARE Tlg1p was unaffected in these circumstances (our unpublished results). Δnvj1 cells in stationary phase showed delocalization of GFP-Osh1p from both of its target organelles (Figure 4D), as expected. These

Figure 3. Osh proteins are spatially distinct. GFP-tagged full-length Osh proteins expressed from the PHO5 promoter in log phase cells. (A) GFP-Osh1p: in many cells a single bright linear structure can be seen adjacent to the vacuole (arrows). Similar localization was also seen with expression at lower levels from the OSH1 promoter (our unpublished results). (B) GFP-Osh2p: in cells with small buds, Osh2p is concentrated in the bud and the neck region of mother cell (arrowhead). It has proved impossible to fix GFP-Osh2p to compare its distribution to plasma membrane and ER markers, but in live cells the fluorescence was coincident with the lipophilic dye FM4-64 when added to cells at 4°C to label the plasma membrane (our unpublished results). However, we cannot formally exclude the possibility that the protein is located to that portion of the endoplasmic reticulum that is adjacent to the plasma membrane. (C) GFP-Osh3p: diffusely cytoplasmic with no discernible subcellular concentration. GFP-tagged Osh1p and Osh3p were expressed from pTL311- and pTL313. GFP-tagged Osh2p was expressed by tagging the ORF in the genome of Δosh1 cells (strain TLY227).
results show that under normal growth conditions Osh1p has a bipartite localization to two separate organelles, one of which is the NV junction, and that by manipulation of growth conditions or genetic background it is possible to target Osh1p solely to one or other of these two locations.

The Ankyrin Repeat Region of Osh1p Localizes to the NV Junction

The dual localization of Osh1p might result either from a single targeting domain binding the same ligand found in two sites, or from two targeting domains each with a distinct ligand specific to a single site. A molecular dissection of Osh1p was undertaken to investigate the basis for its dual localization. Osh1p can be divided into four sections along its primary sequence (Figure 5A). These are a region containing three ankyrin repeats, the PH domain (PH\textsuperscript{Osh1p}), and the oxysterol binding domain (OBD). In addition, there is a less well conserved region between the PH domain and the oxysterol binding domain, which in both Osh1p and OSBP is predicted to contain short stretches of amphipathic helix (Ridgway et al., 1992), and so we will refer to it as the helical domain (HD).

Deletion of the oxysterol binding domain did not affect the bipartite localization seen with full-length Osh1p (Figure 5B), indicating that both targeting signals are found somewhere within the N-terminal three domains. To examine the role of the PH domain in the context of the whole protein, PH\textsuperscript{Osh1p} was replaced with the PH domain of human spectrin, a PH domain that does not show any specific localization when expressed on its own in yeast (our unpublished results). This hybrid construct was no longer localized to punctate structures, but remained localized to the NV junction (Figure 5C), indicating that PH\textsuperscript{Osh1p} is not required for targeting to the NV junction. Subdividing the remaining Osh1p sequences further showed that, while removing the ankyrin repeat region resulted in diffuse fluorescence (PH\textsuperscript{spectrin}.helical domain, Figure 5D), the ankyrin repeats alone localized to the NV junction (Figure 5E). Moreover, a version of Osh1p missing just the ankyrin repeats was found only in small punctate structures, and not the NV junction (Figure 5F). These results demonstrate that the ankyrin repeat domain of Osh1p is necessary and sufficient for targeting of Osh1p to the NV junction.

Formation of the NV junction requires the ER membrane protein Nvj1p, and so we investigated whether the Osh1p ankyrin repeats were binding directly to Nvj1p itself. The NV junction is apparently held together by Nvj1p binding to Vac8p on the vacuolar membrane, so that in \textit{Δvac8} cells the NV junction is lost, and Nvj1p redistributes around the

Figure 4. Osh1p localizes to the NV junction. (A) In log phase, GFP-Osh1p is localized to punctate structures and a single linear structure, lying between the DAPI-stained nucleus and the FM4-64-stained vacuole (arrow). (B) In \textit{Δnvj1} cells, GFP-Osh1p is no longer in linear structures between the nucleus and vacuole. (C) In stationary phase, punctate localization is lost but GFP-Osh1p stays localized to the NV junction. (D) In stationary phase \textit{Δnvj1} cells have lost both localizations, with cytoplasmic staining only. GFP-Osh1p was expressed from plasmid pTL311 in either \textit{Δosh1} cells ("wild type", TLY221), or in \textit{Δnvj1} cells (TLY231). Living cells were labeled with DAPI (nucleus and mitochondria, seen as dots and curvilinear structures) and FM4-64 (vacuoles). The vacuole in the \textit{Δnvj1} cells was often more fragmented than in wild-type cells, an aspect of this deletion not investigated in the original study. It is perhaps not surprising that release from the nucleus should affect the structure of the vacuole, an organelle whose morphology is highly variable depending on growth conditions and strain background (Roberts et al., 1991; Pan et al., 2000).
whole nuclear envelope (Pan et al., 2000). However, in Δvac8 cells the ankyrin repeat region of Osh1p produced entirely diffuse staining (Figure 5G) with no staining of the nuclear envelope where Nvj1p would be expected to be localized (Pan et al., 2000). Vac8p is found in both the NV junction, and on the rest of the vacuolar membrane, but in both wild-type and Δnvj1 neither intact Osh1p nor the ankyrin repeats were seen on the vacuolar membrane outside of the NV junction in the wild-type cells. Taken together, these results indicate that the feature recognized by the ankyrin repeats is neither Vac8p nor Nvj1p alone, but rather a ligand that appears at the NV junction as a result of the interaction of the two proteins.

As shown above, Osh2p does not target to the NV junction, despite also having three ankyrin repeats near its N terminus (Figures 1 and 3). When the N-terminal 285 amino acids of Osh2p, which contain these repeats, was used to replace the equivalent region from Osh1p in the NV-junction–specific construct Osh1DOBDsperpin, it was now diffusely localized, indicating that the ankyrin repeat domains from the two proteins contain different targeting information (Figure 6). Examination of chimeras between the ankyrin repeat domains of the two proteins showed that a region of 100 amino acids of Osh1p, including the third repeat, was sufficient to confer NV-junction targeting activity to the Osh2p chimera (Figure 6). Dividing this region further resulted in loss of targeting, suggesting that the interactions most critical for targeting to the NV junction location are mediated over an extended region within the ankyrin repeat domain that includes repeat 3, and the linker between repeats 2 and 3.

**TGN Targeting Is Specified by the PH Domain of Osh1p**

The above-mentioned results suggest that the PH domain is not required for targeting to the NV junction, but is necessary for targeting of Osh1p to the other punctate structures observed with the full-length protein. These punctate structures have an appearance characteristic of the yeast Golgi, and this was confirmed by with the use of double label immunofluorescence to compare the distribution of the PH-domain containing chimeras with specific compartmental markers. The unstacked nature of Golgi cisternae in *S. cerevisiae* allows different Golgi subcompartments to be distin-

Figure 5. Ankyrin (Ank) repeats of Osh1p specify targeting to the NV junction. (A) Structure of Osh1p and of the constructs examined (plasmids pTL321-325). Osh1p can be divided into four regions: Ank, containing three ankyrin repeats; PH, PHosh1 (in some constructs replaced by PHspectrin); HD, a domain predicted to contain helical regions; and OBD, oxysterol binding domain. The constructs are tagged with GFP ("G"). (B–F) Triple label fluorescent micrographs of live log phase Δosh1 cells (TLY221) expressing the indicated GFP-tagged constructs and stained with FM4-64 (red) and DAPI (blue) as in Figure 4. Arrows indicate typical NV junctions. The same localizations were seen in wild-type cells (our un-published results). The NV junctions are relatively small, and so are not always visible in any given focal plane. When cells expressing GFP-Osh1p were counted the percentage of cells showing clear staining in a linear structure between opposed nuclei and vacuoles was typically 85-95%. The exceptions were almost always small daughter cells where the vacuole is still highly fragmented and presumably the NV junction has yet to properly form. A comparable frequency of NV junction structures was seen with all constructs containing the N-terminal ankyrin repeat domain of Osh1p (G) Δosh1 cells (TLY232) expressing pTL324 treated as in B–F except FM4-64 staining was for 45 min, followed by a 3-h chase to examine transfer of vacuolar material to daughter cells, confirming that loss of Vac8p leads to reduced vacuolar segregation into daughter cells (arrowhead), and highly fragmented vacuoles in some cells (double arrowhead) (Fleckenstein et al., 1998; Pan and Goldfarb, 1998; Wang et al., 1998).
guished by fluorescence microscopy in fixed cells. Although live cells expressing GFP-tagged PH\textsubscript{Osh1} showed punctate localization (Figure 8), this targeting was lost on fixation. In contrast, a C-terminally extended construct consisting of PH\textsubscript{Osh1}+HD was still localized after fixation, although the HD does not add any extra targeting information (Figure 5D), and so this construct was used for colocalization studies. Figure 7A shows that structures containing GFP extensively overlap with Tlg1p, a resident of the TGN and early endosomes (Holthuis et al., 1998). Colocalization between proteins in the Golgi compartments of yeast is usually not absolute, presumably reflecting the dynamic nature of the organelle. In contrast, the distribution of PH\textsubscript{Osh1}+HD was completely distinct both from the cis-Golgi marker Anp1p and the late endosomal marker Pep12p (Figure 7, B and C) (Becherer et al., 1996; Jungmann and Munro, 1998). Thus, the PH domain of Osh1p primarily localizes to TGN membranes or closely related early endosomes. Given that the same construct targeted to the TGN when expressed in mammalian cells (Levine and Munro, 1998), it appears that the nature and localization of the membrane receptor for this family of PH domains has been conserved through evolution.

Figure 6. A portion of the ankyrin repeat domain is required for targeting to the NV junction. Constructs based on Osh1\textsubscript{DOBD}spectrin, a GFP-tagged version of Osh1p without the OBD, and with the PH domain replaced with that of spectrin (plasmid pTPL322, Figure 5C). The ankyrin repeat containing N-terminal region in plasmid pTPL322 was replaced with either the equivalent region of Osh2p, or with the indicated chimeras between Osh1p and Osh2p. The intracellular location of the GFP fusions was examined by fluorescence microscopy of live transfected cells, and the proteins were either entirely diffuse, or localized to the NV junction as indicated. The three ankyrin repeats in Osh1p are 51–80, 96–125, and 196–225 (SWISSPROT P39555).

Figure 7. Immunofluorescence localization of PH\textsubscript{Osh1} domain to the TGN. Δosh1 yeast (TLY221) expressing GFP-tagged PH\textsubscript{Osh1} and the helical domain (residues 280–690, pTL327) were fixed and stained with rabbit antibodies to Tlg1p (A), Anp1p (B), and Pep12p (C), markers for TGN/early endosomes, cis-Golgi, and prevacuolar compartment, respectively. Colocalization of GFP is only seen with Tlg1p (arrowheads), but not with the other two markers (arrows).
targeting of GFP-tagged PH domains was examined. Different intracellular distributions of the full-length proteins are responsible for the different functions, or alternatively the proteins could perform similar functions but be located in different places. We have previously found that the PH domain of mammalian OSBP is the founder member of a protein family with multiple homologues in all eukaryotes so far examined. One possible explanation for such a multiplicity of related genes is that a degree of specialization has arisen on top of a basic function that is common to all. It is possible that such a family of related proteins is located at the same place in the cell, but carries out different functions, or alternatively the proteins could perform similar functions but be located in different places. We have previously found that the PH domain of mammalian OSBP specifies its location to the Golgi, and in this article we investigate the three yeast Osh proteins with PH domains, and find that they have different intracellular distributions. The localizations were determined in live cells expressing GFP-tagged proteins, but because the actual functions of the Osh proteins are still unclear it was not possible to directly examine these functions for each of the GDP fusions. However, for all three proteins it was possible to show that the GFP-fusion was at an accurate representation of the distribution of the native proteins.

PH Domains of Osh Proteins Show Distinct Intracellular Targeting

Full-length Osh1p and Osh2p are localized to different parts of the cell, yet their PH domains are highly homologous (70% identical, 91% similar). To determine whether differences between these PH domains are responsible for the different intracellular distributions of the full-length proteins, targeting of GFP-tagged PH domains was examined. PHOsh1 was clearly visible in punctate structures, while PHOsh2 showed only barely discernible punctate staining, and PHOsh3 was entirely diffuse (Figure 8A). To ask whether this weak membrane localization by PHOsh2 could be enhanced by the presence of surrounding sequences, we also examined constructs comprising the entire N-terminal regions, i.e., the whole protein without the oxysterol binding domains. As mentioned above, Osh1ΔOBD showed a bipartite localization with less in the cytoplasm than seen for the PH domain alone (Figure 8B). However, Osh2ΔOBD surprisingly showed a strong punctate localization similar to PHOsh1, and clearly distinct from the plasma membrane localization of the full-length protein (Figure 8B). This punctate staining by Osh2ΔOBD did not directly result from localization of the regions flanking the PH domain, which when expressed on their own were completely cytoplasmic (our unpublished results). In contrast to the punctate localizations of Osh1ΔOBD and Osh2ΔOBD, Osh3ΔOBD showed partial localization to the plasma membrane. To determine whether these interactions were mediated by the PH domains, conserved basic residues in variable loop 1 of the domain were altered to glutamate residues, mutations which, by analogy with other PH domains, would be expected to abrogate PIP binding (Hyvonen and Saraste, 1997; Levine and Munro, 1998). These mutations caused complete delocalization of PHOsh1, Osh2ΔOBD, and Osh3ΔOBD (Figure 8C), implying that the PH domains of all three Osh proteins can contribute to intracellular targeting. This result indicates that PHOsh2 can target to internal membranes, but the affinity appears weaker than for PHOsh1, and this targeting is apparently masked, or altered, in the context of full-length Osh2p, a situation similar to that seen for mammalian OSBP (Ridgway et al., 1992).

Precise localization of Osh2ΔOBD was not possible, because the construct delocalized under a variety of fixation conditions. However, it would appear to localize to a site closely related to the TGN, because in live cells PHOsh1, Osh1ΔOBD, and Osh2ΔOBD colocalize with FM4-64 at intermediate times after uptake (10–40 min; our unpublished results). It has been shown previously that before accumulating in the vacuole, this endocytic tracer enters the TGN, where it colocalizes with the late Golgi marker Sec7p (Lewis et al., 2000).

DISCUSSION

OSBP is the founder member of a protein family with multiple homologues in all eukaryotes so far examined. One possible explanation for such a multiplicity of related genes is that a degree of specialization has arisen on top of a basic function that is common to all. It is possible that such a family of related proteins is located at the same place in the cell, but carries out different functions, or alternatively the proteins could perform similar functions but be located in different places. We have previously found that the PH domain of mammalian OSBP specifies its location to the Golgi, and in this article we investigate the three yeast Osh proteins with PH domains, and find that they have different intracellular distributions. The localizations were determined in live cells expressing GFP-tagged proteins, but because the actual functions of the Osh proteins are still unclear it was not possible to directly examine these functions for each of the GDP fusions. However, for all three proteins it was possible to show that the GFP-fusion was at least able to rescue phenotypes of deletion strains. Thus, it seems likely that the localizations obtained for these fusions are an accurate representation of the distribution of the native proteins.
We find that Osh1p is located to both the late Golgi and the NV junction. Investigation of the domains required for this dual localization of Osh1p revealed that each intracellular site is specified by a distinct domain in the protein, with the PH domain being necessary and sufficient for Golgi targeting, and the ankyrin repeat domain specifying targeting to the NV junction. This is the first example of a peripheral membrane protein targeting to this specialization of the nucleus and vacuole. A related ankyrin repeat region is also found at the N terminus of some of the OSBP homologues from the yeasts Candida albicans, Kluyveromyces thermotolerans, and S. pombe, but is not present in OSBP from any other organisms, consistent with the notion that the NV junction is a yeast-specific structure. The NV junction is formed by a direct interaction between Nvj1p in the nuclear envelope, and a subpopulation of the Vac8p on the vacuolar membrane (Pan et al., 2000). However, the NV junction-targeting domain from Osh1p does not appear to bind to either of these proteins directly, as revealed by its diffuse distribution in strains lacking either Nvj1p or Vac8p. The targeting thus requires the correct formation of the junctional structure, and so it is possible that the domain binds only to the Nvj1p/Vac8p complex, or to some other factor recruited to the junction. Osh2p has similar ankyrin repeats, but the protein does not localize to the NV junction, and the role of its ankyrin repeats is unclear because this region shows only diffuse localization when fused to GFP (our unpublished results).

In contrast, the PH domain-dependent targeting to the Golgi seems to reflect a more general feature of the OSBP family. We have previously found that the PH domain of mammalian OSBP specifies targeting to the mammalian Golgi (Levine and Munro, 1998), and the same is the case for the Osh1p PH domain in yeast. Indeed this yeast PH domain also specifies Golgi targeting in mammalian cells (Levine and Munro, 1998). The PH domain of mammalian OSBP was found to require PIPs for binding to the Golgi (Levine and Munro, 1998), and such a labile ligand would be consistent with our observation that Osh1p rapidly relocates to the Golgi as cells move out of stationary phase (our unpublished results). A role for PIPs in Golgi targeting is also consistent with phosphatidylinositol 4-kinase having been found on the Golgi in mammalian cells (Godi et al., 2002). A role for PIPs in targeting is also consistent with our observation that Osh1p rapidly relocates to the Golgi (Levine and Munro, 1998). It is, therefore, unlikely that Aur1p is the direct target of the Osh proteins we have

that although Osh2p is normally targeted to the plasma membrane, it has the capacity to translocate to the Golgi in appropriate circumstances.

Intracellular Localization and Function of the OSBP Family

At present the precise function of the OSBP family remains elusive. The necessity and, when overexpressed, the sufficiency of the Osh1p oxysterol binding domain to rescue the phenotype of Δosh1 suggests that the key effects of the protein are mediated by this domain. Although oxysterol binding has so far only been reported for mammalian OSBP, there are several suggestions that yeast OSBP homologues have roles in lipid metabolism and/or trafficking. First, deletion of OSH4 (KE1) is a by-pass suppressor of sec14ts, suggesting a role in regulating phospholipid metabolism in the Golgi (Fang et al., 1996). Second, the growth phenotype of Δosh1 is characteristic of many of the erg mutants of ergosterol synthesis, and has also been reported in disturbances of PIP metabolism (Gaber et al., 1989; Jiang et al., 1994; Stolz et al., 1998). Third, Δosh2, but not Δosh1, is associated with reduced ergosterol levels, although with no growth phenotype (Daum et al., 1999). However the clearest evidence for a role in ergosterol biology comes from a recent comprehensive analysis of the phenotypes of all 127 combinations of OSH gene deletions (Beh et al., 2001). Deletion of all seven OSH genes is lethal, with cells accumulating three-fold elevated levels of ergosterol. Any one OSH gene is sufficient to rescue this lethality (with only OSH1 requiring overexpression), indicating that the seven proteins share a common essential function. However, when each of the OSH genes is deleted individually a specific range of phenotypes is observed, many consistent with mild perturbation of ergosterol synthesis or trafficking, indicating that the encoded proteins perform distinct functions, in addition to their common essential function.

Our localization data provide strong independent support for the idea that the different Osh proteins have specific functions, because it appears that individual proteins are targeted to distinct parts of the cell. A set of proteins acting in a range of different intracellular locations seems to us more compatible with a role in either the sensing of lipid levels, or the transport of lipids or precursors between organelles, rather than with a direct role in lipid synthesis. For instance one possible function is the intracellular traffic of oxysterols, which have been found in a variety of species, including yeast (Nes et al., 1989; Bocking et al., 2000; Gardner et al., 2001). Because oxidation of sterols makes them more water soluble, oxysterols will have a higher propensity to diffuse across the cytoplasm, making them more suitable as second messengers. The presence of high-affinity receptors on the periphery of specific organelles might increase the local concentration of oxysterols present at low levels, and so present a sterol-dependent signal to proteins restricted to the particular bilayer. It has been proposed from work in mammalian cells that OSBP regulates sphingolipid synthesis, which might be expected to be coregulated with the levels of sterols (Lagace et al., 1999). We have recently found that Aur1p, the inositol phosphophorylceramide synthase that initiates yeast sphingolipid synthesis, is localized to the medial Golgi (Levine et al., 2000). It is, therefore, unlikely that Aur1p is the direct target of the Osh proteins we have
studied here, and indeed cells deleted for all three genes showed no change in the activity of Aur1p as measured with the use of a fluorescent substrate in whole cells (our unpublished results). It may be that the effect of OSBP on sphingomyelin synthase is indirect, but it is also possible that it is regulated differently from inositol phosphophorylceramide synthase.

The targeting to the NV junction is intriguing, and because the ER and the vacuole have critical roles in the synthesis and degradation of lipids, respectively, it is conceivable that a structure that connects the two could have a role in the transport of lipid metabolic precursors, or the regulation of lipid metabolism. Interestingly, it has recently been reported that an ER protein Tsc13p, which is required for fatty acid elongation is enriched in the NV junction (Kohlwein et al., 2001). Although the significance of this is somewhat unclear because Elo2p and Elo3p, which are involved in the same process, are not enriched in the junction, it is at least suggestive of a role for the NV junction in lipid metabolism or transport. The NV junction itself is not required for growth on low tryptophan because Δosh1 cells grew as wild type and Δosh1/Δosh1 cells grew as Δosh1, and removal of the ankyrin repeats did not prevent Osh1p rescuing Δosh1 (our unpublished results).

In summary, we have found that 3 OSBP homologues in yeast are spatially distinct, despite all having PH domains, and that Osh1p has a novel targeting determinant specific for the NV junction. This diverse intracellular targeting suggests that the families of OSBP-related proteins present in all eukaryotes are likely to include homologues that perform similar tasks at multiple intracellular sites.

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