The Rab GTPase YPT-1 associates with Golgi cisternae and Spitzenkörper microvesicles in Neurospora crassa

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Summary

Vesicle traffic involves budding, transport, tethering and fusion of vesicles with acceptor membranes. GTP-bound small Rab GTPases interact with the membrane of vesicles, promoting their association with other factors before their subsequent fusion. Filamentous fungi contain at their hyphal apex the Spitzenkörper (Spk), a multivesicular structure to which vesicles concentrate before being redirected to specific cell sites. The regulatory mechanisms ensuring the directionality of the vesicles that travel to the Spk are still unknown. Hence, we analyzed YPT-1, the Neurospora crassa homologue of Saccharomyces cerevisiae Ypt1p (Rab1), which regulates different secretory pathway events. Laser scanning confocal microscopy revealed fluorescently tagged YPT-1 at the Spk and putative Golgi cisternae. Co-expression of YPT-1 and predicted post-Golgi Rab GTPases showed YPT-1 confined to the Spk microvesicular core, while SEC-4 (Rab8) and YPT-31 (Rab11) occupied the Spk macrovesicular peripheral layer, suggesting that trafficking and organization of macro and microvesicles at the Spk are regulated by distinct Rabs. Partial colocalization of YPT-1 with USO-1 (p115) and SEC-7 indicated the additional participation of YPT-1 at early and late Golgi trafficking steps.

Introduction

In eukaryotic cells, the transport of proteins through the distinct membranous compartments of the secretory pathway is carried out in vesicles. The fidelity of the vesicles delivered to the membrane domains of the acceptor compartments requires the coordinated action of GTPases of the Rab family, protein coats, molecular motors, SNARE proteins and tethering factors (Grossshans et al., 2006). Rab proteins are highly conserved small GTPases that regulate vesicle traffic, and have the ability to oscillate between an inactive GDP-bound and an active GTP-bound state. The exchange of the bound GDP for GTP is catalyzed by Guanine nucleotide Exchange Factors (GEFs), while the hydrolysis of GTP to GDP is catalyzed by GTPase Activating Proteins (GAPs). The coordinated actions of GEFs and GAPs on specific Rab GTPases allow them to associate with membrane carriers, and to interact with specific effectors at the acceptor membranes in a regulated manner (Mizuno-Yamasaki et al., 2012).

More than 60 Rabs have been identified in mammalian cells, and about 8 to 12 Ypts (Rab homologues) have been identified in fungi (Pereira-Leal and Seabra, 2000; Schultz et al., 2000; Bourett et al., 2007). In the secretory pathway, the regulation of the anterograde traffic of vesicles has been ascribed mainly to the activity of three Rabs: Rab1, Rab8 and Rab11. In Saccharomyces cerevisiae, Ypt1p, the orthologue of the mammalian Rab1, is involved in membrane traffic from the ER to early Golgi (Jedd et al., 1997; Schultz et al., 2000; Sacher et al., 2001), intra-Golgi (Suvorova et al., 2002) and from early endosomes to late Golgi (Du and Novick, 2001; Lafourcade et al., 2004; Solafani et al., 2010). Ypt31p and Ypt32p are the functional homologues of the mammalian Rab11, involved in the budding of late Golgi vesicular carriers, a traffic event downstream of Ypt1p (Benli et al., 1996; Jedd et al., 1997).

The final stages of the secretory pathway are regulated by the Rab8 orthologue, Sec4p, which has been localized at sites of polarized secretion (Salminen and Novick, 1987) and is implicated in tethering events at the plasma membrane, together with the exocyst complex subunit Sec15p (Guo et al., 1999).

The Golgi apparatus is considered one of the most important sorting organelles for proteins in transit to the plasma membrane. Golgi morphology and organization in
the budding yeast and filamentous fungi differ significantly from the characteristic stacked cisternae described for animal and plant cells (Faso et al., 2009; Pantazopoulou and Peñalva, 2009; Lowe, 2011; Noda and Yoda, 2013). In filamentous fungi, the Golgi appears as individual smooth-surfaced cisternae, fenestrated hollow spheres or tubules (Girbardt, 1970; Grove and Bracker, 1970; Howard, 1981; Pantazopoulou and Peñalva, 2009). It has been suggested that the differences in Golgi organization could correspond in part to the membrane trafficking activity of the Golgi-associated Rab proteins (Liu and Storrie, 2012). Recently, in the filamentous fungus Aspergillus nidulans, alterations in the Golgi organization were linked to RabO, the Ypt1p orthologue (Pinar et al., 2013a). Even though it is widely known that Rab GTPases play significant roles in the anterograde and retrograde traffic of membranes within the secretory pathway in S. cerevisiae or mammalian cells, relatively little is known about their role in hyphal morphogenesis in filamentous fungi.

Fungal hyphae, neurons, root hairs and pollen tubes, all share the unique characteristic of growing at their tips, which results from the asymmetrical and continuous delivery of the necessary precursors for cell wall and membrane expansion (Heath, 1990; Geitmann and Emons, 2000). The origin of the vesicular carriers that transport enzymes responsible for the synthesis of the cell wall components in fungi is presently unclear (Riquelme et al., 2007; Verdín et al., 2009). Most hyphae contain at their apices a specialized multivesicular component, the Spitzenkörper (Spk), which is proposed to act as a Vescicle Supply Center, where cargo-carrying vesicles accumulate before being redirected to tip growing regions (Bartnicki-Garcia et al., 1989). Because of its composition and spatio-temporal localization, the Spk is considered a crucial player in hyphal apical growth and morphogenesis (Riquelme et al., 2011). Evidence from transmission electron microscopy has shown that the main constituents of the Spk are vesicles, actin filaments and ribosomes (Grove and Bracker, 1970; Howard, 1981; Roberson and Vargas, 1994). Two main regions can be defined at the Spk: the ‘core’ or inner region, mainly occupied by microvesicles (20–40 nm in diameter), and the outer region, occupied by macrovesicles (70–100 nm in diameter) (Bartnicki-Garcia, 1990; Riquelme and Sánchez-León, 2014). The microvesicles are indeed chitosomes, which carry chitin synthases (Riquelme et al., 2007; Sánchez-León et al., 2011). At the macrovesicular layer, subunits of the exocyst complex and enzymes necessary for β-(1, 3)-glucan syntheses have been found (Verdin et al., 2009; Riquelme and Sánchez-León, 2014; Riquelme et al., 2014). Little is known about the biogenesis and transport of the cell wall-building vesicular carriers that constitute the Spk. Therefore, in this study we have investigated the intracellular distribution of the Neurospora crassa Rab GTPase YPT-1. Its localization at the Spk core and Golgi cisternae indicates that it participates both in late and early steps of the secretory pathway.

Results

N. crassa YPT-1, YPT-31 and SEC-4 localized at distinct layers of the Spk

A homologue of the small GTPase Ypt1p/Rab1 encoding gene had been previously identified in the N. crassa genome (Heintz et al., 1992; Borkovich et al., 2004). To characterize its subcellular distribution in N. crassa, YPT-1 was fluorescently tagged at its N-terminus and expressed under the control of the glucose repressible promoter ccg-1 (Pccg-1) and the ypt-1 gene promoter (Pypt-1). No differences were observed in the subcellular distribution of GFP-YPT-1 expressed under the control of the Pccg-1 or the Pypt-1 promoters (Fig. 1A, panels 1, 2). Western blots revealed similar expression levels of GFP-YPT-1 when expressed under control of the Pccg-1 or the Pypt-1 promoter (Fig. S1). In both cases, the gfp-tagged version of ypt-1 was directed to the his-3 site, and the selected strains contained the native copy of ypt-1. It was not possible to recover viable homokaryon ypt-1 Δ strains (data not shown). Thus, the functionality of the GFP tagged YPT-1 was not tested. GFP-YPT-1 was observed at the Spk of all vegetative growing hyphae examined (Fig. 1A, panel 1, arrow). To confirm that YPT-1 was indeed confined to the core of the Spk we used the lipophilic dye FM4–64, which has been widely used to stain membranous structures such as the Spk (Fischer-Parton et al., 2000). FM4–64 stained primarily the outer layer of the Spk, surrounding the Golgi-YPT-1 core (Fig. 1B, panel 1). We then further proceeded to analyze whether YPT-1 associates with a specific population of vesicles within the Spk. N. crassa chitosomes are usually observed at the microvesicular region of the Spk (Riquelme et al., 2007; Sánchez-León et al., 2011); therefore, we analyzed the distribution pattern of mChFP-YPT-1 and CHS-1-GFP at the Spk in a heterokaryon strain expressing both recombinant proteins. Co-expression revealed a conspicuous co-localization of both proteins at the Spk core (Fig. 1B, panel 2). Densitometry analysis of the Western blots of fractions obtained by sucrose density gradients (10–65%) showed that although GFP-YPT-1-associated structures (50 kDa bands) were present in all fractions (2–18) analyzed, the highest concentration of GFP-YPT-1 was found within the density range of 1.119–1.138 g ml⁻¹ (fractions 10–12), characteristic of chitosomes (Fig. S2). The N. crassa β-(1, 3)-glucan synthase-related protein (GS-1) has been previously observed at the macrovesicular layer of the Spk (Verdin et al., 2009). The distribution of GS-1 at this layer of the Spk coincides with the distribution of macrovesicles observed...
Fig. 1. Subcellular distribution of Rab GTPases YPT-1, YPT-31 and SEC-4 in Neurospora crassa growing hyphae.

A. LSCM analysis showed accumulation of YPT-1, YPT-31 and SEC-4 at distinct regions of the Spk (arrowheads). 1 and 2: GFP-YPT-1 expressed under the control of a copy of the endogenous promoter (Pypt-1) showed similar distribution patterns as observed in the strain expressing the heterologous protein under the control of the glucose repressible promoter Pccg-1 (McNally and Free, 1988). At the apex, GFP-YPT-1 was observed at the Spk core, whereas at subapical and distal hyphal regions YPT-1 was observed at numerous punctate structures, presumably Golgi cisternae (arrowheads). 3–5: tDIMER-2-YPT-31 and SEC-4 were found at the Spk outer layer (arrowheads). No differences were found in the distribution patterns of SEC-4 expressed under the control of a copy of the corresponding endogenous promoter (Psec-4) or SEC-4 tagged with mChFP expressed under the Pccg-1 promoter. SEC-4 also accumulated at the subapex behind the Spk (arrowhead). 6: GFP-SYN-1 was observed at the macrovesicular and microvesicular layers of the Spk. Bars, 5 μm.

B. 1: Rab GTPases YPT-1, YPT-31 and SEC-4 differentially associate with distinct vesicle populations of the Spk. Labeling with FM4–64 confirmed that GFP-YPT-1 was confined to the Spk core. 2: Co-expression of mChFP-YPT-1 and CHS-1-GFP, a GFP labeled chitin synthase previously found at the microvesicular core of the Spk (Sánchez-León et al., 2011), showed clear co-localization of YPT-1 and CHS-1 at the Spk core. 3: Co-expression of mChFP-YPT-1 and GS-1-GFP, a GFP labeled glucan synthase-related protein identified previously at the macrovesicular layer of the Spk (Verdín et al., 2009), indicated no co-localization of YPT-1 with the macrovesicular outer layer of the Spk. 4: YPT-31 was localized at the Spk outer layer surrounding the YPT-1 core. 5: The fluorescently tagged Rabs YPT-31 and SEC-4 were localized at the Spk outer layer. Note the variable labeling pattern of YPT-31 and SEC-4 at the outer vesicular layer of the Spk. 6: YPT-31 and the GFP tagged exocyst subunit EXO-70 (Riquelme and Sánchez-León, 2014; Riquelme et al., 2014) showing partial co-localization at the Spk macrovesicular area. Bars, 2 μm.
by transmission electron microscopy of hyphal tips (Grove and Bracker, 1970; Howard, 1981). To further confirm that YPT-1 is excluded from the macrovesicular region of the Spk, we analyzed heterokaryon strains co-expressing GS-1-GFP and mChFP-YPT-1. It was clearly observed that mChFP-YPT-1 and GS-1-GFP occupied distinct layers of the Spk (Fig. 1B, panel 3).

In *S. cerevisiae* Rabs, Ypt31/32p associate with the late Golgi and newly formed post-Golgi carriers, where they participate in the recruitment of Sec2p, the GEF for Sec4p (Benli *et al*., 1996; Jedd *et al*., 1997; Ortiz *et al*., 2002). Sec4p participates downstream of Ypt31/32p at the last traffic events of the exocytic vesicles (Salminen and Novick, 1987; Goud *et al*., 1988). A previous analysis identified *N. crassa* YPT-31 (NCU01523) and SEC-4 (NCU06404) as the putative orthologues of the *S. cerevisiae* post-Golgi Rabs Ypt31/32p and Sec4p respectively (Borkovich *et al*., 2004). Blast search alignment of the *N. crassa* putative Rabs (SEC-4 and YPT-31) showed high overall identities (62–87%) at the amino acid level with their corresponding fungal and human orthologues (Table S2, Fig. S3). To determine whether YPT-31 and/or SEC-4 were involved in the traffic of vesicles that arrive at the Spk in *N. crassa*, we fluorescently tagged both Rabs at their N-terminus with the red fluorescent protein tdIMER-2 and GFP or mCherry, respectively, while maintaining their corresponding native copies. Confocal microscopy analysis of tdIMER-2-YPT-31 revealed its accumulation at the macrovesicular outer layer of the Spk (Fig. 1A, panel 3, arrow). Co-expression of tdIMER-2-YPT-31 with GFP-YPT-1 in a heterokaryon strain clearly confirmed the distribution of each protein in different layers of the Spk (Fig. 1B, panel 4). SEC-4 was mainly found at the outer macrovesicular region of the Spk (Fig. 1A, panel 4, arrow), but also some cytoplasmic fluorescence could be found at subapical hyphal regions (Fig. 1A, panel 4, arrowhead). Expression of the fluorescently tagged SEC-4 under the control of *Pccg-1* or the *Psec-4* promoters revealed no differences at the subcellular distribution (Fig. 1A, panels 4 and 5). Co-expression of tdIMER-2-YPT-31 and GFP-SEC-4, allowed the identification of both Rabs at the macrovesicular layer of the Spk, although the co-localization was only partial (Fig. 1B, panel 5). The distribution pattern of the Rabs YPT-31 and SEC-4 at the hyphal tips was variable due to the pleomorphic characteristics of the *N. crassa* Spk. To determine the accumulation of secretory vesicles at the Spk, the putative v-SNARE protein SYN-1 (NCU00566.7), used in other fungal species as vesicular marker (Lewis *et al*., 2000; Taheri-Talesh *et al*., 2008), was tagged with GFP at its N terminus. vSNAREs are members of the synaptobrevin protein family and are involved in fusion events of exocytic carriers during late steps of the secretory pathway (Protopopov *et al*., 1993; Grote *et al*., 2000; Shanks *et al*., 2012). A Blast search alignment revealed that *N. crassa* SYN-1 has a high identity level with the vSNAREs SynA of *A. nidulans* (64%; NP_014972) and Snc2p of *S. cerevisiae* (58%; NP_014972). GFP-SYN-1 was observed at the macrovesicular and microvesicular layers of the Spk (Fig. 1A, panel 6; Fig. S4). In *N. crassa*, the components of the exocyst complex, EXO-70, EXO-84 and SEC-3, were found associated to the periphery of the Spk (Riquelme and Sánchez-León, 2014; Riquelme *et al*., 2014). We co-expressed tdIMER-2-YPT-31 and the previously GFP-tagged exocyst subunit EXO-70 (expressed under control of its own promoter), and observed partial co-localization of YPT-31 and EXO-70 at the Spk macrovesicular layer, although EXO-70 occupied primarily the area between the plasma membrane and YPT-31 labeled area (Fig. 1B, panel 6).

**Vesicles flow into the Spk at very high rates**

To determine the turnover rates of microvesicles and macrovesicles, we conducted fluorescence recovery after photobleaching (FRAP) analysis of GFP-YPT-1 or tdIMER-2-YPT-31. Photobleaching of the fluorescence at the Spk lasted 5 s. No fluorescence signal was detected at the Spk area immediately after photobleaching (Fig. 2A, panel 1–4, dotted circle at time 0.0 s). Fluorescence recovery of GFP-YPT-1, FM4–64, CHS-1-GFP, tdIMER-2-YPT-31 and GS-1-GFP occurred progressively at the Spk (Fig. 2A), where fluorescent spots appeared a few seconds after photobleaching (Fig. 2A, arrows). An increasing amount of GFP-YPT-1 or CHS-1-GFP over time was detected at the microvesicular/chitosomal core of the Spk, whereas the accumulation of FM4–64 occurred in the region surrounding the area occupied by GFP-YPT-1 (Fig. 2A, panels 1 and 2). Fluorescence recovery of tdIMER-2-YPT-31 and GS-1 occurred also progressively, but restricted to the macrovesicular layer of the Spk (Fig. 2A, panels 3 and 4). Half-time recovery values (*t*½, Mean ± Standard Error), i.e. the time at which half of the final fluorescence intensity was reached, were assessed from the normalized fluorescence recovery values for GFP-YPT-1, FM4–64, tdIMER-2-YPT-31, CHS-1-GFP and GS-1-GFP (Fig. 2B). The *t*½ was 15.5 ± 4.3 s (*n* = 11) for GFP-YPT-1, 21.3 ± 5.7 s (*n* = 11) for FM4–64, 21.5 ± 3.7 s (*n* = 9) for CHS-1-GFP and 9.9 ± 2.6 s (*n* = 9) for tdIMER-2-YPT-31 and 10.2 ± 1.7 s (*n* = 12) for GS-1-GFP. The one-way analysis of variance (ANOVA) post hoc Tukey-HSD test revealed significant differences between the *t*½ of tdIMER-2-YPT-31 and GS-1-GFP compared with the *t*½ of CHS-1-GFP and FM4–64, whereas no significant differences were found between the *t*½ of GFP-YPT-1 and the rest of *t*½ analyzed.

Total internal reflection fluorescence microscopy (TIRFM) analysis confirmed the conspicuous localization
mChFP-YPT-1 at vesicles (Fig. 3A and B, arrowheads) and putative Golgi cisternae (Fig. 3A and B, arrows) moving fast and slow respectively. Kymographs confirmed not only variable speeds of the YPT-1 labeled structures (Fig. 3C, panel 1), but also that the movements were primarily parallel to the axis of hyphal growth (Fig. 3C, panel 2). Traces obtained from the kymographs revealed vesicles with fast (0.4–4 μm s⁻¹, n > 50) anterograde and retrograde movements (Fig. 3C, panels 3 and 4), which contrasted with the slow (< 0.5 μm s⁻¹, n > 20) anterograde motion recorded for the putative Golgi cisternae (Fig. 3C, panel 5).

YPT-1 associates with diverse subpopulations of Golgi cisternae

In addition to the Spk localization, GFP-YPT-1 was also observed as numerous fluorescent punctate structures along the cell (Fig. 1A, panel 1, arrowheads), which were distributed mainly at subapical and distal hyphal regions (8–49 μm from the tip, n = 28), although some labeled particles were observed at the proximal subapical hyphal regions near the tip (7 μm from the tip, n = 28). To determine if the GFP-YPT-1 punctate structures corresponded to Golgi cisternae, we examined the effects of brefeldin A..
(BFA) on the intracellular distribution of GFP-YPT-1. The addition of BFA to growing hyphae induced the formation of GFP-YPT-1 clusters and a reduction of putative Golgi punctate structures (Fig. 4A, Movie S1). When exposed to BFA, accumulation of GFP-YPT-1, tDIMER-2-YPT-31 and GFP-SEC-4 at the hyphal apex was temporarily interrupted, concomitant with a temporary cessation of apical growth, tips swelling and Spk disassembly of the Rabs (Fig. 4A). BFA treatment of hyphae co-expressing CHS-1-mChFP or tDIMER-2-YPT-31 and GFP-YPT-1 showed clustering of the YPT-1 punctate structures, but no accumulation of CHS-1 or YPT-31 at any of these structures (Fig. S5, Movie S1). Interestingly, during BFA exposure we observed CHS-1-mChFP at the distinct subpopulations of Golgi cisternae, we co-expressed GFP- or mChFP-YPT-1 with different Golgi markers such as USO-1, VRG-4, VPS-52 and SEC-7, and exposed the cells to BFA. In \( S. \) \( \text{cerevisiae} \) and mammals, Uso1p/p115 is a long coiled-coiled tethering protein, which is recruited by Ypt1p/Rab1 (Cao \textit{et al.}, 1998; Allan \textit{et al.}, 2000) and is necessary, through the interaction with GM130 (Barr \textit{et al.}, 1998; Shorter and Warren, 1999; Moyer \textit{et al.}, 2001) for the tethering of ER-derived membrane carriers to early Golgi membranes (Beard \textit{et al.}, 2005). The expression of USO-1-GFP under the control of its own promoter showed numerous fluorescent punctate structures distributed along the hyphae (Fig. 4B). The morphology and subcellular distribution of the USO-1-GFP labeled structures resembled that of the putative Golgi cisternae marked by YPT-1 (Fig. 1A, panel 1). At distal and subapical distal hyphal regions (> 36.2 \( \mu \)m from the tip, \( n = 30 \)), USO-1-GFP and mChFP-YPT-1 co-localized at numerous particles, presumably early Golgi cisternae (Fig. 4C, panel 1, arrowheads; Fig. 4D). Although both proteins co-localized at some structures, the pattern of the fluorescence signal exhibited pleomorphisms (Fig. 4C, panel 1, 3–6; Movie S2), which might result from the dynamic behavior of the Golgi cisternae during maturation. Numerous mChFP-YPT-1 structures that did not co-localize with USO-1-GFP were also observed along different hyphal regions (Fig. 4C, arrows). After BFA exposure, a change in the distribution of fluorescent cisternae was clearly observed (Fig. 4C, panel 2). Visual inspection of the fluorescent signals suggested that almost all USO-1-GFP labeled cisternae co-localized with a fraction of the mChFP-YPT-1 labeled cisternae (Fig. 4C, panel 2, bottom image). To verify the extent of co-localization, the Intensity Correlation Quotient (ICQ) was calculated from the images corresponding to hyphae treated with BFA and untreated. The ICQ index measures the relationship of the relative intensities of two fluorescent particles (Li \textit{et al.}, 2004). A random distribution of two fluorescent particles has an ICQ value around 0, while co-localizing and nonco-localizing particles have ICQ values within the range of 0 < ICQ < +0.5 and 0 > ICQ ≥ −0.5 respectively. In untreated cells, ICQ values (Mean ± Standard Error) were 0.144 ± 0.026 (\( n = 8 \)), whereas in BFA-treated cells ICQ values were significantly higher, 0.212 ± 0.019 (\( n = 5 \)) (Student’s t-test, \( P < 0.01 \)).

The Vrg4 protein is a mannose transporter, whose localization has been reported at Golgi structures in different fungal systems (Dean \textit{et al.}, 1997; Nishikawa \textit{et al.}, 2002; Jackson-Hayes \textit{et al.}, 2008; Bowman \textit{et al.}, 2012). Experimental evidence on the GDP-mannose transporter of \( S. \) \( \text{cerevisiae} \), Vrg4p, suggests that the protein recycles between ER and Golgi (Abe \textit{et al.}, 2004). To further analyze the distribution of YPT-1 at the different Golgi cisternae populations, we co-expressed mChFP-YPT-1 and VRG-4-GFP. Image analysis revealed co-localization of VRG-4-GFP and mChFP-YPT-1 in a subpopulation of Golgi cisternae (Fig. 4E, panel 1, arrowheads; Movie S3). Although co-localizing structures were present, numerous single labeled structures were observed (Fig. 4E, panel 1, arrows). BFA exposure induced the typical clustering of mChFP-YPT-1 labeled cisternae, which co-localized with all the VRG-4-GFP labeled cisternae (Fig. 4E, panel 2, arrowheads). The ICQ value increased from 0.091 ± 0.025 (\( n = 12 \)) in untreated cells to 0.201 ± 0.025 (\( n = 10 \)), significantly higher (Student’s t-test, \( P < 0.01 \)), in BFA-treated cells.

To determine if the YPT-1 single labeled structures observed in the above experiments corresponded to late Golgi cisternae, co-localization analyses were performed with the \( N. \) \( \text{crassa} \) putative orthologue of Sec7p and YPT-1. In \( S. \) \( \text{cerevisiae} \), \( \text{SEC7} \) encodes a guanine-nucleotide exchange factor, which activates Arf GTPases at the late (trans-) Golgi cisternae (Achstetter \textit{et al.}, 1988; Sata \textit{et al.}, 1998). An analysis of the intracellular localization and dynamics of Golgi in \( S. \) \( \text{cerevisiae} \) and \( A. \) \( \text{nidulans} \) revealed that Sec7 could be used as a bona fide marker of late Golgi cisternae (Loise \textit{et al.}, 2006; Matsuura-Tokita \textit{et al.}, 2006; Pantazopoulos and Peñalva, 2009). Blast search alignment revealed that the \( N. \) \( \text{crassa} \) NCU07658.7 encodes a protein with an overall identity of 38% and 57% with the \( S. \) \( \text{cerevisiae} \) Sec7p (NP_010454) and \( A. \) \( \text{nidulans} \) HrpB\textsuperscript{Sec7} (AN6709) proteins respectively. Additionally, the predicted amino acid sequence has the Sec7 domain (Pfam: PF01369). Therefore, the \( N. \) \( \text{crassa} \) NCU07658.7 was designated as Sec7. The endogenously GFP tagged SEC7 was identified at several fluorescent punctate structures, presumably late Golgi cisternae, distributed along the cells (Fig. 5A, \( n > 30 \)); occasionally, some of these structures were observed near hyphal tips (Fig. 5A, arrowheads). As predicted, the co-expression of SEC7-GFP and mChFP-YPT-1 revealed that both proteins co-localized at some of the putative late Golgi cisternae (Fig. 5B, panel 1). This co-localization increased slightly
**Fig. 4.** YPT-1 associates with different subpopulations of Golgi cisternae.

A. BFA treatment in cells expressing the fluorescently tagged YPT-1. YPT-31 and SEC-4 did not cause any clustering of tDIMER-2-YPT-31 or GFP-SEC-4 as observed for GFP-YPT-1. The Spk localization of the Rabs was temporarily interrupted while the hyphae experienced tip swelling. Bars, 5 μm.

B. LSCM analysis showed USO-1-GFP at numerous punctate structures distributed at the proximal and distal subapical hyphal regions. Bars, 5 μm.

C. 1: Co-expression of mChFP-YPT-1 and the early Golgi marker USO-1-GFP (arrowheads) showed co-localization of YPT-1 and USO-1 at some of the Golgi cisternae located at distal and subapical distal regions of the cells. 2: BFA exposure caused the aggregation of USO-1-GFP and mChFP-YPT-1 (arrowheads). Note that a small fraction of mChFP-YPT-1 clusters (arrows) did not co-localize with USO-1-GFP. 3–6: Digital magnification of co-localizing particles in panel 1 (arrowheads) shows both proteins occupying distinct domains within the identified Golgi cisternae. Bars, 5 μm.

D. Illustration of the subcellular distribution of cisternae containing mChFP-YPT-1 and USO-1-GFP and their positions from the tip (Mean ± Standard Error).

E. 1: Co-expression of mChFP-YPT-1 and VRG-4-GFP (Bowman *et al.*, 2012) showed a mixed population of co-localizing (arrowheads) and single (arrows) labeled cisternae. 2: mChFP-YPT-1 clusters resulting from the BFA exposure co-localized with VRG-4-GFP containing cisternae (arrowheads). Bars, 5 μm.
after BFA treatment (Fig. 5B, panel 2). ICQ values before and after BFA treatment were $0.122 \pm 0.037$ ($n = 5$) and $0.154 \pm 0.030$ ($n = 8$) respectively. Unexpectedly, BFA treatment promoted the accumulation of SEC-7-GFP at the Spk (Fig. 5B, panel 2; $n > 20$), in contrast to the untreated cells.

To further analyze whether the YPT-1 labeled structures corresponded to late Golgi compartments, intracellular distribution of YPT-1 and the GARP complex subunit VPS-52 was analyzed. The GARP (Golgi-associated retrograde protein) complex is a multimeric tethering factor involved in the retrograde traffic of endosome-derived carriers to late Golgi or TGN (Conibear and Stevens, 2000). The GARP complex, composed of VPS-51p, -52p, -53p and -54p, is recruited by Ypt6p/Rab6 to the late Golgi (Siniossoglou and Pelham, 2001; Liewen et al., 2005). In A. nidulans, the yeast Ypt6p homologue, RabC(Rab6), was identified at early and late Golgi compartments (Pantazopoulou and Peñalva, 2011). In previous studies, localization analysis in N. crassa identified the fluorescently tagged VPS-52 at Golgi cisternae mainly separated from the early Golgi cisternae labeled by the calcium transporter PMR-1, which lead the authors to suggest that VPS-52 localizes at late Golgi cisternae (Bowman et al., 2012). Previous studies in S. cerevisiae found Ypt1p at late Golgi cisternae, and reported that this Rab GTPase regulates the traffic of vesicles at the early endosome-late Golgi interface (Sclafani et al., 2010). To determine if a subpopulation of Golgi cisternae labeled with YPT-1 corresponded to late Golgi cisternae, we co-expressed tDIMER-2-VPS-52 and GFP-YPT-1. As predicted, some of the tDIMER-2-VPS-52 containing Golgi cisternae coincided with GFP-YPT-1 Golgi cisternae (Fig. 5C, panel 1, arrowheads). Image analysis yielded an ICQ of $0.177 \pm 0.013$ ($n = 4$). Under the effects of BFA, tDIMER-2-VPS-52 was dispersed into the cytosol (there was an increase in the red
fluorescence signal of the cytoplasm) and was no longer observed in the putative late Golgi cisternae (Fig. 5C, panel 2). Co-expression of SEC-7-GFP and tDIMER-2-VPS-52 showed both proteins partially co-localizing at the same structures (Fig. 5D, panel 1 and 3). Co-localization analysis yielded an ICQ of 0.113 ± 0.017 (n = 21). BFA treatment induced the aggregation of SEC-7-GFP marked Golgi and the dispersion of tDIMER-2-VPS-52 to the cytosol (Fig. 5D, panel 2).

YPT-1 distributes along a tubular network

In addition to the identification of YPT-1 at Golgi cisternae, and at the core of the Spk (see results above), we found YPT-1 at other subcellular membranes. At distal hyphal regions (79.6 ± 22.8 μm, n = 7), mChFP-YPT-1 was found decorating numerous tubule-like structures (Fig. S6A, panel 1). None of these tubular structures were observed near the distal subapical or apical regions of the hyphae. TIRFM analysis showed that the length and position of the YPT-1 tubules varied along the hyphae (Fig. S6B). The tubular structures frequently arranged into a loose network of filaments that displayed a very dynamic behavior (Fig. S6C; Movie S4); sometimes, they were found forming branches and loops (Fig. S6C). To examine the possibility that the YPT-1 tubules corresponded to microtubules, we co-expressed mChFP-YPT-1 and the GFP tagged β-tubulin (BML-GFP). No co-localization of any of the structures was found, confirming that the YPT-1 tubule-like structures are not associated with microtubules (Fig. S6A, panel 2–4; Movie S5).

YPT-1 does not localize at early endosomes

Previous studies have shown that Rab1/Ypt1p has a role in endocytic traffic (Sclafani et al., 2010; Mukhopadhyay et al., 2011). While several of our observations indicated that YPT-1 localized primarily at the different Golgi cisternae, we could not exclude the possibility that some of the GFP- or mChFP-YPT-1 labeled structures corresponded to early endosomes. Moreover, partial co-localization was observed between very few of the GFP-YPT-1 cytoplasmic punctate fluorescent structures and FM4–64 stained structures (Fig. S7A). To discern this, we co-expressed mChFP-YPT-1 with the previously GFP-tagged YPT-52, the N. crassa early endosome Rab5 homologue (Seidel et al., 2013). None of the GFP-YPT-52 labeled endosomes co-localized with mChFP-YPT-1 (Fig. S7B; Movie S6). Each of the samples analyzed (n = 8) showed ICQ values of –0.5.

Discussion

The Spk is a pleomorphic structure implicated in the hyphal growth and morphogenesis of numerous fungal species (Riquelme and Sánchez-León, 2014). Previous studies have revealed that the conspicuous Spk of N. crassa hyphae contains a high concentration of vesicles of different sizes organized in different layers, and more importantly, that each type of vesicle contains cell wall-building enzymes of different nature (Riquelme et al., 2007; Verdin et al., 2009). Previously, RabC and RabO, the proteins in A. nidulans homologous to Rab6 and Rab1, respectively, were observed at the Spk (Pantazopoulou and Peñalva, 2011; Pinar et al., 2013a). In this study, we identified the Rab proteins YPT-1, YPT-31 and SEC-4 at different layers of the Spk of N. crassa (Fig. 6A), suggesting that distinct Rabs might differentially regulate the traffic of the different populations of vesicles of the Spk. Fluorescent tagging of some Rab GTPases in other model systems has resulted in hypomorphic effects due to alterations in function and/or

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stability of the tagged proteins (Pantazopoulou et al., 2014). We expressed fluorescently tagged versions of YPT-1, SEC-4 and YPT-31, while maintaining their corresponding native copies, resulting in strains with no growth defects (data not shown).

The steady fluorescence at the Spk suggests that the rate of discharge of vesicles from the Spk equals the rate of arrival of the vesicles to the Spk, consistent with the vesicle supply center (VSC) model for fungal morphogenesis, which proposed that secretory vesicles arriving from distal regions of the hyphae accumulate temporarily at the Spk prior to being discharged to the apical plasma membrane (Bartnicki-Garcia et al., 1989) (Fig. 6B). The turnover rates obtained from analysis of FRAP experiments indicated that the putative secretory vesicles exhibit a highly dynamic behavior at the Spk (Jones and Sudbery, 2010; Dijksterhuis and Molenaar, 2013; Pantazopoulou et al., 2014). FRAP analysis in N. crassa revealed a half-time recovery (t½) for YPT-1 and YPT-31 between 10 and 16 s, suggesting the existence of high flow rates for vesicles reaching the Spk. In A. nidulans and C. albicans, a t½ of 7 s and 10 s was obtained in FRAP experiments for the YPT-31 orthologue, RabERAB11 and Sec4 respectively (Jones and Sudbery, 2010; Pantazopoulou et al., 2014). In spite of the differences in growth rates reported for hyphae of N. crassa (∼ 60 μm min⁻¹) (Hickey et al., 2002), C. albicans (0.25 μm min⁻¹) (Jones and Sudbery, 2010) and A. nidulans (0.5–1 μm min⁻¹) (Horio and Oakley, 2005), similar t½ values were obtained for the above-mentioned Rabs, indicating that the rate of arrival of secretory carriers to the Spk is independent of hyphal growth rate. The t½ values obtained for YPT-1 and YPT-31 were lower than those for FM4–64 and CHS-1, which could be indicative of the transient association of the Rabs with the secretory carriers, in contrast with CHS-1 and FM4–64, which are presumably embedded in the plasma membrane of the carriers. These results hint at the existence of a mechanism coordinated by Rabs that regulates the ordered delivery of vesicles to the Spk and their subsequent discharge to the apical cell surface.

We identified the N. crassa YPT-1 at the core of the Spk, co-localizing almost fully with CHS-1 (Fig. 6A). In a recent study, we have identified YPT-1 among the proteins that co-immunoprecipitated with CHS-1, CHS-4 and CHS-5 (Fajardo-Somera, unpublished). In addition, we found that GFP–YPT-1 sedimented in fractions with a density range similar to the density (1.125 g ml⁻¹) of the fractions with high chitin synthase activity reported previously (Bartnicki-Garcia et al., 1984; Leal-Morales et al., 1994; Verdín et al., 2009). Altogether, the evidence suggests that YPT-1 might have a role in the pre-exocytic traffic of microvesicles, including chitosomes. The putative role of YPT-1 in regulating vesicle traffic at the Spk, adds yet another role for Ypt1, other than the role described in S. cerevisiae and mammalian cells at the ER-Golgi interface, endosome-Golgi and autophagosome formation (Barrowman et al., 2010; Lynch-Day et al., 2010). This functional ‘promiscuity’ of Ypt1 at different steps of the secretory pathway must be precisely regulated by corresponding GEFs and GAPs.

Ypt31/32p have been localized at sites of polar growth in N. crassa (Jedd et al., 1997). They participate in a cascade mechanism, where they recruit to the membrane of secretory carriers Sec2p, a GEF that subsequently activates Sec4p (Ortiz et al., 2002). An analysis of the amino acid sequence of N. crassa Rab GTPases SEC-4 and YPT-31 revealed a high level of identity with Sec4p/Rab8 and Ypt32p/Rab11 respectively. In addition, sequence alignments showed a remarkable level of sequence conservation at the identified regions (G motifs, Switch Regions I and II, and RabSF1–4). Together, these results suggest that SEC-4 and YPT-31 of N. crassa are indeed homologous proteins of Rab8 and Rab11 respectively. In N. crassa, we found YPT-31 at the peripheral layer of the Spk surrounding the YPT-1 core (Fig. 6A). Given the distribution of both FM4–64 and YPT-31 at the peripheral layer of the Spk, we expected to obtain similar rates of recovery from bleaching for the fluorescent probe and the Rab protein. However, the t½ of YPT-31 was significantly lower than the t½ of FM4–64, suggesting that YPT-31 associates with a distinct subpopulation of macrovesicles at the Spk. An alternative explanation would be that FM4–64 labels membrane carriers that arrive at the Spk from a separate secretory route derived from endosomal recycling. Although additional biochemical approaches are needed to decipher the differential association of YPT-1 and YPT-31 with the vesicles of the Spk, the results of the localization experiments and the differences in t½ indicate that the stratification of the Spk is delicately choreographed by Rab GTPases. Furthermore, we identified that SEC-4 distributed similarly to YPT-31 at the outer layer of the Spk (Fig. 6A). These results are consistent with the suggested role of Ypt31/32p and Sec4p in later traffic steps of the secretory pathway. Sec4c has been previously identified at hyphal tips of C. albicans, Aspergillus fumigatus and Ashbya gossypii (Schmitz et al., 2006; Jones and Sudbery, 2010; Powers-Fletcher et al., 2013). Notably, in N. crassa, not only SEC-4 but also YPT-31 is involved in the traffic of Spk macrovesicles. Recently, the characterization of the N. crassa exocyst complex revealed that subunits SEC-3, EXO-70 and EXO-84 partially associate with the outer macrovesicular layer of the Spk (Riquelme and Sánchez-León, 2014; Riquelme et al., 2014). In C. albicans, based on the differences in the dynamics and localization patterns observed for Sec4 and exocyst components Exo70 and Exo84, it has been proposed that secretory vesicles of the Spk associate with the exocyst at the cell surface (Jones and Sudbery, 2010). In N. crassa,
the partial co-localization of EXO-70 and YPT-31 at the Spk outer layer, suggests that the membrane carriers labeled by YPT-31 associate with the exocyst subunits (i.e. EXO-70) at the periphery of the Spk before being directed to exocytic sites. It remains to be tested whether any of these exocyst subunits serves as effectors of SEC-4 or YPT-31.

It is well recognized that the Golgi apparatus is an organelle that has a pivotal role in the secretory system of eukaryotic cells (Mowbrey and Dacks, 2009). In contrast to the stacked organization of the Golgi apparatus seen in some organisms, the fungal Golgi cisternae (early and late compartments) are scattered and optically resolvable by light microscopy (Wooding and Pelham, 1998). According to the proposed model for Golgi cisternal maturation, secretory cargoes are modified by a specific set of enzymes, which change over time within the same compartment (cisterna) (Losev et al., 2006; Matsuura-Tokita et al., 2006). This model assumes that the enzymes utilized in the early Golgi are recycled back from late Golgi through membrane carriers traveling in a retrograde manner; accordingly, a late Golgi cisterna corresponds to a mature early Golgi cisterna that has acquired ‘late proteins’. Within the diverse fungal groups, there is wide evidence of variation in the organization and morphology of the Golgi apparatus. For instance, in contrast to the stacked cisternae of Golgi observed in Schizosaccharomyces pombe and Pichia pastoris, N. cerevisiae, N. crassa and A. nidulans have dispersed Golgi cisternae (Mogelsvang et al., 2003; Pantazopoulou and Peñalva, 2005; Suda and Nakano, 2012; Pinar et al., 2013b). Protein secretion processes at Golgi require the orchestrated actions of distinct Rabs to ensure vesicle directionality to specific organelles (Hutagalung and Novick, 2011). The S. cerevisiae Ypt1p and the mammalian Rab1 have been implicated in the regulation of the traffic of vesicles from and to the distinct Golgi structures (Barrowman et al., 2010). In addition to the localization at hyphal tips, we identified YPT-1 at numerous punctate structures along the hyphae in N. crassa. The distribution pattern, pleomorphic nature and motility of these structures were similar to the structures previously labeled by various Golgi markers (Hubbard and Kaminskyj, 2008; Bowman et al., 2012; Pinar et al., 2013a). Our co-localization analysis of YPT-1 and putative early and late Golgi markers confirmed that this Rab GTPase associates with Golgi cisternae and participates in traffic events at the Golgi interface in N. crassa. A recent study in A. nidulans showed that BFA affects the morphology of the Golgi by promoting the aggregation of its cisternae (Pinar et al., 2013a). In N. crassa, exposure to BFA induced the formation of clusters of Golgi cisternae, which were labeled by GFP-YPT-1, but not with tDIMER-2-YPT-31, confirming that YPT-1 participates at the membrane trafficking events at the Golgi level. The lack of YPT-31 at similar structures and its exclusive localization at hyphal tips suggest that the N. crassa YPT-31 participates in post-Golgi steps of the secretory pathway, as previously shown for YPT-31/32 in the budding yeast (Jedd et al., 1997) and for RabERab11 in A. nidulans (Pantazopoulou et al., 2014). Under exposure to BFA, arrival of YPT-1 and YPT-31 to the Spk was only transiently blocked, while Golgi cisternae organization was disturbed. This suggests that YPT-1 and YPT-31 associated vesicles either reach the Spk by a Golgi-independent secretory pathway, as previously suggested for CHS (Riquelme et al., 2007), or alternatively, that the gradual recovery of Golgi cisternae from BFA effects allows some exit of secretory carriers destined to the Spk. Recently, the Ypt1p orthologue in A. nidulans, RabO

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in traffic steps between early endosomes and late Golgi. YPT-1 could be acting on the target membrane in endosome-to-Golgi traffic. Further mutant and biochemical analyses are needed to further understand their mode of operation.

Thus far, little is known about the relationship of Rab GTPases and the traffic of vesicles of the Spk. A better understanding on the role of Rab GTPases at late secretory traffic events would help to decipher the organizational differences found in the Spk of diverse filamentous fungal species.

**Experimental procedures**

**Strains and culture conditions**

Bacterial and _N. crassa_ strains used or generated in this study are listed in Table S1. _Escherichia coli_ DH5α was grown on LB medium (1% tryptone, 0.5% yeast extract, 1% NaCl) supplemented with ampicillin (100 μg ml⁻¹) and incubated at 37°C. _N. crassa_ cells were routinely grown in Vogel’s minimal medium (VMM) containing 1% sucrose and solidified with 1.5% agar when needed (Vogel, 1956). For auxotrophic His strains, histidine (0.5 mg ml⁻¹) was added to VMM medium. Transformed conidia were plated on VMM-FGS (0.5% fructose, 0.5% glucose, 20% sorbose) supplemented with hygromycin B (200 μg ml⁻¹; InvivoGen) when required. For crosses, mat a conidia were spread over _mat A_ mycelium grown on solid synthetic crossing medium with 2% sucrose (Westergaard and Mitchel, 1947). For growth analysis, elongation rate of _N. crassa_ transformants was measured on VMM agar plates at 30°C.

**Recombinant DNA techniques and plasmid constructions**

Plasmids and oligonucleotides used are described in Table S1. Standard polymerase chain reaction (PCR) and cloning procedures were used to fuse the _sgfp_, _mcherry_ or _tdimer2_ gene to the 5’ end of the ORFs of the Rab GTPases _ypt-1_ (NCU08477.7), _ypt-31_ (NCU01523.5), _sec-4_ (NCU06404.7) or the vSNARE _syn-1_ (NCU00566.7) in the plasmids PCCG::N-GFP (Honda and Selker, 2009), pJV18-N and pMF334 (Freitag and Selker, 2005) respectively. The open reading frames (ORFs) of _ypt-1_ (1067 bp), _ypt-31_ (837 bp), _sec-4_ (959 bp) and _syn-1_ (980 bp) were identified in _N. crassa_ (www.broadinstitute.org/annotation/genome/neurospora/MultiHome.html) and amplified by PCR from _N. crassa_ N1 (FGSC988) genomic DNA with custom-designed primers that included restriction endonucleases sites at the 5’ and 3’ termini. PCR was performed in a Bio-Rad Thermal Cycler using TaKaRa LA Taq® DNA polymerase (Clontech). PCR was performed with the following conditions: denaturation at 94°C (2 min) was followed by 30 cycles of 94°C (30 s), 55°C (30 s) and 72°C (2–2.5 min), and a final extension at 72°C (5 min). For the _sgfp_ gene fusion, the amplified and gel purified PCR products of the _ypt-1_ and _sec-4_ genes were digested with Ascl and XbaI and inserted into Ascl/XbaI-digested PCCG::N-GFP (GeneBank accession No. FJ457006) yielding plasmids.
pESL07b-2 and pESL05b-2; while the PCR product of syn-1 was digested with PacI and XbaI, and inserted in PCCG::N-GFP plasmid digested with the same restriction endonucleases. For the tdimer2 gene fusion, the amplified and gel purified PCR product of the ypt-31 gene was digested with MluI and BglII and inserted into MluI/BglII-digested pMF334 (GeneBank accession No. DQ250999.1), yielding plasmid pdimer2(12)-YPT-31. For the expression of ypt-1 and sec-4 under the control of their endogenous promoters, we amplified ~1 kb of the 5’ untranslated region of the corresponding ORFs, and inserted the PCR digested products into EcoRI/BamHI digested pCoS204 (Seidel et al., 2013), replacing the cccg-1 promoter (Pcccg-1). Amplified and gel purified PCR ORFs of ypt-1 and sec-4 were PacI/XbaI digested and subcloned in the corresponding promoter containing vectors.

To obtain mcherry gene fusions, the PCR products of ypt-1 and sec-4 were digested with PacI and EcoRI and inserted into PacI/EcoRI digested pJV18-N yielding plasmids pESL10-25 and pESL08-15 respectively. For the sgp tagging of putative early and late Golgi markers, we searched the N. crassa genome sequences for orthologues of the S. cerevisiae coiled-coil tethering protein USO-1 (NCU01644.7) and SEC-7 (NCU07658.7), respectively, and designed primers to tag uso-1 and sec-7 endogenously at their C-terminus by using the split marker gene replacement method (Smith et al., 2011). The primers (Table S1) design and PCR strategy for the above technique were previously described (Riquelme and Sánchez-León, 2014; Riquelme et al., 2014). Additionally, we designed the primer pairs USO-1-F/USO-1-R-Gly and SEC-7-F-01/SEC-7-10xGLY-R to amplify ~1 kb fragments of the uso-1 and sec-7 ORFs, respectively, immediately upstream of the stop codon; while the primer pairs USO-1-UTR-F-lox/USO-1-UTR-R and LOX-3UTR-SEC-7-F/3UTR-SEC-7-R were designed to amplify ~1 kb fragments of the uso-1 and sec-7, of their 3’ untranslated corresponding region. The resulting fused PCR fragments were gel purified, mixed in equal proportions (~0.5 µg DNA per transformation), and used to transform N. crassa NMF263 (FGSC9718) and/or N2928 (FGSC9717) conidia.

Neurospora genetics

Conidia of N. crassa host strains FGSC9717 or FGSC9718, deficient in nonhomologous end-joining (Ninomiya et al., 2004), were transformed with either Ndel or SspI linearized vectors or PCR products by electroporation in a Bio-Rad Gene Pulser (capacitance, 25 µF; voltage, 1.5 kV; resistance, 600 Ω) as previously described (Riquelme et al., 2007). For each transformation, 20 to 30 histidine protrophs (His⁺) or hygromycin B (200 µg ml⁻¹) resistant (Hyg⁺) transformants were transferred to VMM slants and screened for the expression of GFP, mChFP or tdimer2 by confocal microscopy. Crossing of transformants was performed as previously described (Riquelme et al., 2007). To select homokaryons, the resulting ascospores were inoculated onto VMM plates, heat shocked at 60°C for 1 h and incubated at 25°C. No viable homokaryon knockouts strains could be recovered for any of the Rabs analyzed, which hindered testing the functionality of any of the fluorescently tagged Rabs by complementation.

Blast search for N. crassa proteins SEC-4, YPT-31, SYN-1 and SEC-7 was performed using the GeneBank accession numbers XP_962242.1, XP_956972.2, XP_964727.2 and XP_962785.2 respectively (Altschul et al., 1990). Alignments of the Rab GTPases with their corresponding orthologues were performed in Geneious Basic 4.8.5 software (using the default program settings). Sequences of fungal and human Rabs identified with high level of identity were obtained using the GeneBank accession numbers listed in Table S2.

Microscopy analysis and image processing

Imaging of the transformant strains was conducted in three different confocal microscopes:

i. An inverted Zeiss LSM-510 Meta laser scanning confocal microscope (LSCM) fitted with an argon/2 ion laser (GFP: excitation, 488 nm; emission, 500 to 550 nm); For simultaneous visualization of GFP and mCherry an argon/2 and He/Ne-2 ion laser (mCherry: excitation 543–561 nm; emission, 575 to 615 nm) along with a 100X Plan Neofluar oil-immersion objective (NA, 1.3) or 63X LC Plan Neofluar Imm Korr DIC (NA, 1.3) were used for the confocal observations. Confocal images were captured using LSM-510 software (version 3.2; Carl Zeiss) and evaluated with a Carl Zeiss LSM Image Examiner software (version 4.2).

ii. An inverted Olympus Fluoview™ FV1000 confocal microscope fitted with an argon laser (GFP: excitation, 488 nm; emission, 505–525 nm) and a diode pump solid state (DPSS, Melles Griot, Carlbad, CA) laser (mCherry or RFP: excitation, 543 nm; emission, 560–660 nm) was used for individual or sequential visualization of GFP and mCherry or tdimer2. A 60X Plan Apo N (Olympus) oil-immersion objective (NA, 1.42) was used with this equipment. Confocal images were captured and examined using FV10-ASW software (version 4.0.2.9, Olympus).

iii. An upright Leica DM5000 confocal microscope fitted with an argon laser (GFP: excitation, 488 nm; emission, 497–536 nm) and a DPSS laser (mCherry or tdimer2: excitation, 561 nm; emission, 568–646 nm) were used for individual or simultaneous visualizations of the fluorescent molecules. A 63X HCX PL APO lambda blue oil-immersion objective (NA, 1.4) was used. Confocal images were captured using TCS SP5 Leica System and examined using LAS AF software (version 2.6.3. Leica). For FRAP analysis, the fluorescent Spk of growing hyphae was photobleached for 5 s with the 488 nm argon laser or with 543 nm DPSS laser at ~50% of intensity, and fluorescence recovery images were captured at 0.61 to 3.39 s intervals at 2.5–5% of laser intensity. Fluorescence signal intensities of the Spk were analyzed with ImageJ (version 1.48u; Maryland, USA). Time series of images were converted to grayscale and the average pixel intensity of the Spk area was measured at every time point. Intensity values were normalized, considering the highest average intensity value after photobleaching as the 100%. Half-time recovery time (t1/2, Mean ± Standard Error), i.e. the time at which half of the final fluorescence intensity was reached was calculated from the average of t1/2 obtained from individual samples. For TIRFM, an IX-81 inverted microscope equipped with a 60X (NA, 1.49) Apo-N TIRF objective lens and DPSS lasers: 488 nm and 561 nm (Melles Griot, Carlbad, CA), was used. Images were recorded with a Hamamatsu Digital Camera C11440 Orca-Flash 2.8 (Hamamatsu Photonics K.K., Japan) for durations of 2–3 min at 960 × 720
pixels resolution and 0.9 s per frame. Dimensions cell Sens software (version 1.9; Olympus Corporation) was used to control the camera and capture images. All images were further processed with Adobe Photoshop CS5 (version 15.0). Kymographs and the moving rates of YPT-1 fluorescent particles were obtained from TIRFM time series (70 frames in 64 s; 120.9 nm/pixel) using the Ykymomaker software (Chiba et al., 2014); the automatic trace detection function of the software was used to calculate the moving rates of individual particles that exhibited only short distance movements (∼2.3 μm). One-way ANOVA, the post-hoc Tukey HSD test and Student’s t-test statistical analysis were performed on the STATISTICA data analysis software system (version 8.0, StatSoft, Inc.). For time-lapse and live-cell imaging, the inverted agar block method was employed (Hickey et al., 2005). For microscopy analyses of the hyphae under BFA treatment, a thin coverslip (0.17 μm, 22 × 22 mm) was placed between the microscope slide and the inverted agar block containing the growing hyphae (∼0.5 mm away from the edge of the colony). After scanning untreated hyphae, the coverslip was subtly lifted to directly add 10 μl of BFA so we could immediately monitor the inhibitory effects over the same hyphae in less than 1 min. To analyze the in vivo co-localization of two proteins, heterokaryons were obtained on VMM agar plates, where conidia of two different N. crassa strains of the same mating type were co-inoculated and the ICQ was assessed by Western blot analysis of the GFP-YPT-1 constructs driven by the Ppyt-1 promoters, fresh conidia from the corresponding strains were grown in Vogel’s liquid medium at 30°C and 150 r.p.m. for ∼20 h. Mycelia were harvested by filtration, washed twice with sterile cold deionized water, and pulverized with liquid nitrogen. Pulverized mycelia (5–8 ml) were mixed with 25 ml of lysis buffer (Tris-HCl 50 mM, pH 7.4; KCl 100 mM; MgCl2 10 mM; Triton X-100 1%) and protease inhibitors. Disrupted cells were centrifuged at 2850 × g (Rmax) for 20 min, and the supernatant was centrifuged at 19 000 × g at 4°C for 1 h. Two hundred microliters of the supernatant of each sample was mixed with Laemmli buffer (2×), boiled for 5 min and kept at −20°C until their analysis by Western blot.

For Western blot analysis, 30 μl of the selected fractions or 40 μg of total protein equivalent as bovine serum albumin (BSA) were separated by 8–13.5% SDS-PAGE. Resolved proteins were transferred to Nitrocellulose (Pall Corporation) or PVDF (Millipore) filter membranes, which were immunoreacted for GFP- or mChFP-tagged proteins with a mouse anti-GFP antibody 1:1000 (Santa Cruz Biotechnology, CA) and a peroxidase-conjugated anti-mouse antibody 1:3000 (Pierce, Rockford, IL). Membranes were incubated first with a 1:1000 anti-BFA antibody and then with the appropriate secondary antibody. The blots were immersed in Pierce enhanced chemiluminescent reagent (Pierce, Rockford, IL) and exposed to X-ray film. All protein bands were quantified with ImageJ software (version 1.48u; Maryland, USA). For microscopy analyses of the hyphae under BFA treatment, a thin coverslip (0.17 μm, 22 × 22 mm) was placed between the microscope slide and the inverted agar block containing the growing hyphae (∼0.5 mm away from the edge of the colony). After scanning untreated hyphae, the coverslip was subtly lifted to directly add 10 μl of BFA so we could immediately monitor the inhibitory effects over the same hyphae in less than 1 min. To analyze the in vivo co-localization of two proteins, heterokaryons were obtained on VMM agar plates, where conidia of two different N. crassa strains of the same mating type were co-inoculated and grown overnight at 28–30°C. For quantitative correlation image analysis of co-localization, the ICQ was assessed using ImageJ software (version 1.48u; Maryland, USA). Fluorescence intensities of the hyphal regions (∼60 μm) analyzed were adjusted and the obtained values normalized.

Fluorescent dye and inhibitors

A stock solution (20 mg ml−1) of the fungal metabolite brefeldin A (BFA; Sigma) was prepared in dimethyl sulfoxide and prepared fresh at a final concentration of 200–300 μg ml−1 in VMM. For staining of the endocytic membranes, we used 10–25 μM of the styryl dye FM–64 (N-(3-triethylammoniumpropyl)-4-(6-(4-(diethylamino)phenyl)hexatrienyl)pyridinium dibromide) (Molecular Probes®). For each dye or inhibitor, the agar blocks containing hyphae were inverted onto coverslips containing 5 to 10 μl of the diluted working stock.

Density gradient centrifugation and Western blots

Conidia (1 × 104 cells ml−1) from the selected transformed strains were inoculated into 200–500 ml Vogel’s complete medium (VCM) and incubated at 30°C, under 220 r.p.m. agitation, for 20 h. Mycelia were harvested by filtration, washed twice with sterile cold deionized water, and once with homogenization buffer (33 mM phosphate buffer, pH 8.2). Mycelia (∼10 g) were mixed with 10 ml homogenization buffer (33 mM, pH 8.2), protease inhibitors, 20 g of glass beads (425–600 microns, Sigma), and were disrupted at a low temperature in a Braun MSK homogenizer for 4 × 30 s. Disrupted cells were centrifuged at 1000 × g (Rmax) at 4°C for 10 min, and the supernatant was layered on top of a 35 ml linear sucrose gradient (10–65%, w/v) in homogenization buffer. The mix was centrifuged at 184 000 × g (Rmax) for 4 h, at 4°C in a Beckman ultracentrifuge using a 70Ti rotor. The gradient was fractionated from the top with an ISCO fractionator using 70% (w/v) sucrose as changing solution. Fifty to 100 μl of each fraction were mixed with Laemmli buffer (2×), boiled for 5 min and kept at −20°C until their analysis by Western blot. For expression analysis of the GFP-YPT-1 constructs driven by the Ppyt-1 and Ppyt-1 promoters, fresh conidia from the corresponding strains were grown in Vogel’s liquid medium at 30°C and 150 r.p.m. for ∼20 h. Mycelia were harvested by filtration, washed twice with sterile cold deionized water, and pulverized with liquid nitrogen. Pulverized mycelia (5–8 ml) were mixed with 25 ml of lysis buffer (Tris-HCl 50 mM, pH 7.4; KCl 100 mM; MgCl2 10 mM; Triton X-100 1%) and protease inhibitors. Disrupted cells were centrifuged at 2850 × g (Rmax) at 4°C for 25 min, and the supernatant was centrifuged at 19 000 × g at 4°C for 1 h. Two hundred microliters of the supernatant of each sample was mixed with Laemmli buffer (2×), boiled for 5 min and kept at −20°C until their analysis by Western blot.

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**Supporting information**

Additional supporting information may be found in the online version of this article at the publisher’s web-site.