**FolVps9, a Guanine Nucleotide Exchange Factor for FolVps21, Is Essential for Fungal Development and Pathogenicity in *Fusarium oxysporum* f. sp. lycopersici**

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The soil-borne, asexual fungus *Fusarium oxysporum* f.sp. *lycopersici* (*Fol*) is the causal agent of tomato wilt disease. Autophagy plays a crucial role in the development and virulence of *Fol*. The *Fol* endosomal system is highly dynamic and has been shown to be associated with conidiogenesis and pathogenicity. Rab GTPases and the regulators are highly conserved in regulating autophagy and endocytosis in most eukaryotes. Identification and characterization of additional Rab regulators in fungal pathogens should facilitate the understanding of the autophagy and endocytosis in different filamentous fungi. Here, we have identified and characterized the yeast VPS9 homolog *FolVPS9* in *Fol*. Targeted gene deletion showed that *FolVPS9* is important for growth, conidiation and virulence in *Fol*. Cytological examination revealed that *FolVps9* co-localized with *FolVps21* (a marker of early endosome) and played a critical role in endocytosis and autophagosome degradation. Pull-down assays showed that *FolVps9* interacted with *FolVps21*, which was also important for development and plant infection in *Fol*. Yeast two-hybrid, bimolecular fluorescence complementation and co-immunoprecipitation assays revealed that *FolVps9* specifically interacts with the GDP-bound form of *FolVps21*. Furthermore, a constitutively active form of *FolVps21* (Q72L) was able to rescue defects of *ΔFolvps9* and *ΔFolvps21* mutants. In summary, our study provides solid evidence that *FolVps9* acts as a *FolVps21* guanine nucleotide exchange factor (GEFs) to modulate endocytosis and autophagy, thereby controlling vegetative growth, asexual development and pathogenicity in *Fol*.

**Keywords:** *Fusarium oxysporum*, *FolVps9*, guanine nucleotide exchange factor (GEF), plant infection, endocytosis, autophagy

**INTRODUCTION**

Endocytosis is a conserved transport process in which macromolecules, lipids, or membrane proteins are transported to endosomal compartments. Recently, a series of studies have shown that endocytosis plays an important role in cell polarity, signal transduction and plant infection in phytopathogens (Wedlich-Soldner et al., 2000; Ramanujam et al., 2013; Li L. et al., 2017;
During endocytosis, cargo is packaged by an area of plasma membrane, which then buds off inside the cell to form transport vesicles (Balderhaar and Ungermann, 2013; Li X. et al., 2017). These transport vesicles transporting the cargo from inside the cell membrane to the early endosome undergo a sorting process, in which they are either trafficked back to the plasma membrane or sent to the vacuole (Seaman, 2008; Li X. et al., 2017; Zhu et al., 2018).

Previous studies have shown that Rab GTPases regulate endocytic vesicle trafficking by targeting these vesicles to appropriate cellular compartments (Balderhaar and Ungermann, 2013; Ramanujam et al., 2013; Zheng et al., 2015). For example, the early stage of the endocytic pathway is regulated by Rab5, while Rab7 controls the late stage (Balderhaar and Ungermann, 2013; Solinger and Spang, 2013). Rab proteins exist in both the active GTP-bound and inactive GDP-bound states, which are regulated by guanine nucleotide exchange factor (GEFs) and GTPase-activating proteins (GAPs) (Zerial and McBride, 2001; Markgraf et al., 2007; Zheng et al., 2018a). GEFs facilitate formation of the GTP-bound form of the Rab protein; in this state, the Rab protein can bind multiple tethering effectors to promote vesicle trafficking (Ishida et al., 2016). Conversely, GAPs catalyze hydrolysis of GTP to GDP on the Rab protein, thereby converting it to the GDP-bound conformation (Cai et al., 2007; Ishida et al., 2016). Based on sequence similarity and their cognate GTP-binding protein, GEFs have been divided into different families (Cherfils and Chardin, 1999; Hutagalung and Novick, 2011; Ishida et al., 2016). To date, more than 40 putative Rab GEFs have been identified in humans, and they all possess at least three conserved domains, including the Vps9, DENN, and Sec2 motifs (Ishida et al., 2016).

In Saccharomyces cerevisiae, there are at least three Vps9 domain proteins (Vps9p, Muk1p, and Vrl1p) that function as Rab5 GEFs (Bean et al., 2015). Vps9p was the first GEF identified, and it has been shown to bind retromer (an endosome-localized complex involved in protein retrograde) to facilitate recruitment of PI3P (phosphatidyl-inositol-3-phosphate) to endosomes (Bean et al., 2015). Subsequent studies have shown that Vps9p is a vacuolar sorting protein that contains a Vps9 domain, which is involved in vesicle-mediated vacuolar protein trafficking (Hama et al., 1999). Previous studies have shown that Vps9p, Muk1p and Vrl1p have redundant, yet distinct functions (Bean et al., 2015). Deletion of MUK1 does not result in a visible phenotype in yeast. However, the VPS99 MUK1 double mutant displays a more severe phenotype than the VPS9 single mutant (Paulsel et al., 2013). In addition, overexpression of MUK1 or VRL1 fully restores the temperature-sensitive growth phenotype of Δvps9, but cannot replace the role of Vps9 in late endosomal sorting of carboxypeptidase Y (CPY; Paulsel et al., 2013; Bean et al., 2015).

Cross-talk between autophagy and endocytosis exists and there is evidence demonstrating that the same proteins play important roles in both processes (Lipatova et al., 2012; Sarkar, 2013). However, the endocytosis pathway is mainly responsible for absorption and utilization of nutrients from outside the cell, while the autophagy pathway is mainly responsible for the degradation of endogenous waste components from the cytoplasm. During autophagy, cellular components are packaged by coated vesicles to form autophagosomes, which then fuse with the vacuole to facilitate degradation (Voigt and Poggeler, 2013). In addition to autophagy-related proteins (Atgs), other proteins that regulate vesicle trafficking, including Rab and its GEFs, are essential for autophagy (Gao et al., 2013; Bean et al., 2015; Li et al., 2015; Liu et al., 2015; Zheng et al., 2015). In S. cerevisiae, the Rab5p/Vps21p module controls formation of autophagosomes and Rab7 regulates fusion of autophagosomes with the vacuole (Chen et al., 2014; Zhou et al., 2017).
objective (Zeiss, Oberkochen, Germany). Liquid CM was used to grow cultures for extraction of genomic DNA and total RNA.

**Fungal Transformation and Gene Deletion**

*F. oxysporum* protoplast preparation and fungal transformation were performed as previously described (Li et al., 2019a). The split-marker PCR approach was used to generate targeted gene replacement vectors (Catlett et al., 2003; Fang et al., 2018; Yang et al., 2018). Primers used for amplifying flanking sequences for each gene are listed in Supplementary Table S2. These primers were used to amplify fragments for transformation using PCR, after which the PCR products were purified and transformed into protoplasts of wild type strain Fol4287. Hygromycin B (Solarbio, Beijing, China) was added to agar medium at a final concentration of 300 μg ml⁻¹ for transformant selection. The putative targeted gene deletion mutants were screened by PCR assays with two pairs of primers in Supplementary Table S2 and further confirmed by Southern blots.

**Vector Construction**

Unless noted otherwise, all fluorescent protein fusion vectors were constructed with the entire protein coding sequence under the control of RP27 constitutive promoter. Fragments were amplified and transformed with Xhol-digested pYF11 into *S. cerevisiae* strain XK1-25 and final vector assembled using the yeast gap repair approach. For the FolVps21 domain deletion and the S27N, Q72L point mutation vectors, primers were designed to use in Splicing Overlap Extension (SOE)-PCR to create the mutations, amplifying sequences from the RFP-FolVPS21 fusion vector. The PCR products were cloned into the pYF11 plasmid using yeast gap repair and verified by sequencing analysis. The primers used in this section are listed in Supplementary Table S2.

**Virulence Assays**

For tomato root inoculation assays, 12 days old seedlings (Moneymaker, susceptible i-2) were carefully uprooted and washed in tap water to remove soil particles. The tomato seedling roots were inoculated using a conidial suspension (5 × 10⁶ conidia ml⁻¹) or mock (water) for 30 min. The treated seedlings were then planted in vermiculite and maintained in a growth chamber. Disease symptoms were examined and photographed 3 weeks after inoculation. Pathogenicity experiments were performed as previously described (Li et al., 2019a). The resulting mixture was centrifuged at 4°C for 30 min at 20,000 × g. The supernatant was incubated with anti-FLAG M2 agarose beads (Sigma-Aldrich) at 4°C for 6 h, and washed three times with wash buffer [50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.5 mM EDTA]. The proteins were eluted with 200 mM glycine (pH 2.5) and then digested with trypsin and analyzed by Liquid Chromatography-Mass Spectrometry/Mass Spectrometry (LC-MS/MS) as described (Li B. et al., 2017). The resulting MS data were analyzed and searched against the non-redundant *F. oxysporum* f.sp. *lycopersici* protein database at NCBI.

**Yeast Two-Hybrid (Y2H), Co-immunoprecipitation (Co-IP) and Bimolecular Fluorescence Complementation (BiFC) Assays**

The yeast two-hybrid assay was carried out using the Matchmaker Gal4 two-hybrid system 3 (Clontech, United States). Wild type FolVps21-AD, FolVps21S27N-AD and FolVps21Q72L-AD were cloned into pGADT7 (prey construct), while wild type FolVps9- BD, FolVps9Δ Vps9-BD and FolVps9Δ GUR-BD were cloned into pGBK7 (bait construct). The pairs of prey and bait constructs were co-transformed into the yeast strain AH109 as previously described (Li B. et al., 2017). pGADT7-T and pGBK7-T-53 were used as positive controls and pGADT7-T and pGBK7-lam were negative controls. The Trp⁺ and Leu⁺ yeast transformants were isolated and assayed for growth on SD dropout medium lacking tryptophan, leucine, histidine and adenine (SD-Trp-Leu-His-Ade).

For co-immunoprecipitation assays, the FolVps9 ORF was cloned into pHZ126 to generate the 3 × FLAG fusion construct and the FolVps21S27N ORF was cloned into pYF11 to generate the GFP fusion construct using the yeast gap repair approach. The resulting constructs were verified by sequencing analysis and subsequently co-transformed into protoplasts of Fol4287. Hygromycin B and G418 (Solarbio, Beijing, China) were added at a final concentration of 300 μg ml⁻¹ for transformant selection. The transformants were confirmed by western blot assays using anti-GFP or anti-FLAG antibody (Abcam, Cambridge, MA, United States). Total proteins were extracted and incubated with GFP beads (Sigma-Aldrich) as previously described (Li L. et al., 2017). Proteins eluted from beads were analyzed by western blot with anti-GFP (GFP-Tag Mouse mAb, Abmart, China) or anti-FLAG antibody (Abcam, Cambridge, MA, United States). Results were visualized with the ECL detection system (Amersham Biosciences).

For BiFC assays, the FolVps9-YFP C and FolVps21S27N-YFP N fusion constructs were made by cloning fragments containing the FolVps9 and FolVps21S27N ORFs along with their native promoter regions (1500 bp upstream from the start codon) into pHZ65 or pHZ68, respectively. The resulting constructs were verified by sequencing analysis and co-transformed into protoplasts of Fol4287. Hygromycin B and G418 (Solarbio,
Gene sequences are available using the following accession numbers: FolVPS9 (FOXG_00470), FolVPS21 (FOXG_09392).

Staining and Cellular Localization Assays
To examine endocytosis, hyphae of the indicated strains were cultured in liquid CM medium at 28°C for 16 h and stained with FM4-64 following procedures previously described (Fischer-Parton et al., 2000). Photographs were taken using a confocal laser scanning microscope (Zeiss, Oberkochen, Germany). For vacuole staining, CMAC (7-amino-4-chloromethylcoumarin; Sigma-Aldrich; stock solution in 10 mM dimethyl sulfoxide) was used to stain hyphae cultured as described above (Zhang et al., 2016).

Autophagy Assays
For GFP-FolAtg8 and cleaved GFP assays, the GFP-FoLATG8 vector was transformed into the Fol4287 wild type strain and various mutants using the polyethylene glycol (PEG)-mediated transformation as previously described. Transformants expressing GFP-FolAtg8 were grown in CM medium for 24 h and then shifted to nitrogen starvation (MM-N) for 8 h to induce non-selective autophagy. Hyphae induced for 8 h were analyzed for GFP fluorescence using a fluorescent microscope. GFP-FolAtg8 proteolysis was assessed by western blot using GFP antibody (GFP-Tag Mouse mAb, Abmart, China). The amount of free GFP and GFP-FolAtg8 was calculated using grayscale analysis (Image-pro plus; Media Cybernetics, Shanghai, China) after western blotting.

Quantitative Real-Time PCR (qRT-PCR) Analysis
Total RNA was extracted from tomato roots using the TRIzol LS reagent (Invitrogen) according to the manufacturer’s recommendations, and subsequently further purified using RNAeasy Mini spin columns (Qiagen). Synthesis of first-strand cDNA was performed as previously described (Li et al., 2019a). The relative quantification of each transcript was calculated using the 2−ΔΔCt method (Livak and Schmittgen, 2001). All qRT-PCR assays were conducted in triplicate for each sample and the experiment was repeated three times. All primers used in this section are listed in Supplementary Table S2.

Statistical Analysis
Each result was presented as the mean ± standard deviation (SD) of at least three replicated measurements. Significant differences between treatments were statistically evaluated by SD and one-way analysis of variance (ANOVA) using SPSS 2.0. The data for two different treatments were compared statistically by ANOVA, followed by the F-test if the ANOVA result was significant at p < 0.01.

Accession Number
Gene sequences are available using the following accession numbers: FolVPS9 (FOXG_00470), FolVPS21 (FOXG_09392).

RESULTS
Identification and Deletion of FolVPS9 in F. oxysporum f. sp. lycopersici
Using the S. cerevisiae Vps9p protein sequence as the reference to blast the F. oxysporum f. sp. lycopersici genome database, we identified FOXG_00470 as a homolog of S. cerevisiae Vps9p that we named FolVPS9. FolVPS9 is predicted to contain 2548 base pairs with two introns, and encode a polypeptide of 783 amino acids (aa). Domain analysis revealed that FolVps9 possesses Vps9 (480-595 aa) and ubiquitin-binding domains (CUE, 741–782 aa)2. To explore the physiological function of Vps9 in F. oxysporum f. sp. lycopersici, we used a gene replacement construct to mutate FolVPS9 (Supplementary Figure S1A). The △Folvps9 mutant was confirmed by Southern blot analysis (Supplementary Figure S1B). The mutation was also complemented by re-introducing the full length FolVPS9 gene (△Folvps9/FolVPS9 strain) and showing that this construct restored wild type functions.

Folvps9 Is Important for Vegetative Growth and Asexual Reproduction
To explore the influence of FolVPS9 on vegetative growth, we first checked the growth and colony morphology of the △Folvps9 mutant on CM and PDA agar plates. After 5 days of incubation in the dark at 28°C, the △Folvps9 mutant exhibited significantly reduced radial growth on CM and PDA media (Figure 1A). The colony diameter was reduced by 53.9 and 44.1% on CM and PDA plates, respectively (Table 1). This result indicates that FolVps9 is important for normal vegetative growth.

Conidia play a key role in infection and subsequent development of pathogenic fungi in the host root (Mes et al., 1999; Li et al., 2019b). Therefore, we investigated the role of FolVps9 in conidia formation. Wild-type, △Folvps9 and △Folvps9/FolVPS9 strains were observed following the induction of conidiophores and conidia formation on carboxymethyl cellulose (CMC) media for 5 days. Microscopic observations showed that conidiophore production by the △Folvps9 mutant was reduced over 50.0% compared with wild type and complemented strains (Figure 1B). Furthermore, compared to wild type, Microconidia was decreased over 60.0% in the △Folvps9 mutant and the conidia produced were smaller (Figure 1C and Table 1). In addition, the macroconidia were reduced 50% in the in the △Folvps9 mutant (Table 1). These results suggest that FolVps9 is important for vegetative growth and conidiogenesis in Fol.

Folvps9 Is Required for Fol Virulence in Tomato Roots
To assess whether FolVps9 is required for pathogenesis of Fol, infection assays were performed using roots from the susceptible cultivar Moneymaker and conidia from wild type, △Folvps9 and △Folvps9/FolVPS9 strains. At 21 dpi, plants inoculated with △Folvps9 showed no obvious disease symptoms and were...
as healthy as the water-treated (mock) plants (Figure 2A). In contrast, plants treated with the wild type and ΔFolvps9/FolVPS9 strains exhibited obvious disease symptoms, with severe stunted growth and wilting (Figure 2A).

To gain further insight into the defect of the ΔFolvps9 mutant on disease progression in tomato, the vascular bundles of plant stems inoculated with conidia of wild type, ΔFolvps9 and ΔFolvps9/FolVPS9 strains were assessed using stereomicroscope at 14 dpi. The results showed that the stem vascular bundles of plants inoculated with wild type and ΔFolvps9/FolVPS9 strains exhibited brown pigmentation, suggesting that the vascular bundles contain lesions, while the color of vascular bundles in the plants infected with the ΔFolvps9 mutant were similar to those of water-treated control plants (Figure 2B).

Fungal recovery assays were used to further assess the pathogenesis defect of the ΔFolvps9 mutant. Compared to the wild type and ΔFolvps9/FolVPS9 strains, infected stems from the ΔFolvps9 mutant yielded very little fungal growth (Figure 2C). These observations were supported by qPCR fungal biomass quantification results, with significantly higher levels of fungal DNA in the wild type and complemented strains than in the ΔFolvps9 mutant (Supplementary Figure S2). Taken together, these results indicated that FolVps9 was required for plant infection in Fol.

**FolVps9 Is Localized in Endosomes and Functions During Endocytosis**

To further understand the role of FolVps9, we checked its subcellular localization. We constructed an expression cassette for GFP-FolVps9 driven by a strong constitutive ribosomal protein promoter, RP27, and transformed into Fol. As mentioned above, this construct rescued the phenotypic defects of the ΔFolvps9 mutant. Fluorescent microscopy of vegetative hyphae from this strain showed that GFP-FolVps9 was localized in punctate cytoplasmic bodies (Figure 3A). Because in yeast Vps9p acts as a GEF for Vps21p, a protein involved in vesicle trafficking and a marker protein for early endosomes, we speculated that the punctate bodies in this strain showed that GFP-FolVps9 was localized in endosomes.

**TABLE 1 | Characterization of growth and conidiation in F. oxysporum f. sp. lycopersici strains.**

| Strain | Colony diameter (cm) | Microconidia | Macroconidia | Microconidia length |
|--------|----------------------|--------------|--------------|---------------------|
| WT     | 6.3 ± 0.2            | 3.1 ± 0.2    | 0.6 ± 0.1    | 9.5 ± 2.1           |
| ΔFolvps9 | 3.4 ± 0.3*          | 1.2 ± 0.1*   | 0.3 ± 0.1*   | 4.8 ± 1.8*          |
| ΔFolvps21 | 2.7 ± 0.2*         | 1.5 ± 0.2*   | 0.5 ± 0.2    | 5.1 ± 2.3*          |
| ΔFolvps9/FolVPS9<sup>PVPS9</sup> | NA         | 1.0 ± 0.3*   | 0.2 ± 0.1*   | NA                  |
| ΔFolvps9/FolVPS9<sup>DCUE</sup> | NA         | 3.1 ± 0.2    | 0.5 ± 0.2    | NA                  |
| ΔFolvps9/FolVPS21<sup>S27N</sup> | NA         | 1.2 ± 0.2    | 0.2 ± 0.1*   | 4.5 ± 2.1*          |
| ΔFolvps9/FolVPS21<sup>Q72L</sup> | NA         | 3.0 ± 0.3*   | 0.3 ± 0.2*   | 9.5 ± 2.1           |
| ΔFolvps9/FolVPS21 | NA       | 1.3 ± 0.2*   | 0.3 ± 0.1*   | 4.7 ± 1.9*          |
| ΔFolvps9/FolVPS21<sup>S27N</sup> | NA         | 1.6 ± 0.1*   | 0.2 ± 0.1*   | 4.9 ± 2.5*          |
| ΔFolvps9/FolVPS21<sup>Q72L</sup> | NA         | 3.1 ± 0.2    | 0.5 ± 0.2    | 9.3 ± 2.0           |
| ΔFolvps9/FolVPS21 | 6.1 ± 0.2   | 3.2 ± 0.1    | 0.6 ± 0.2    | 9.3 ± 0.8           |
| ΔFolvps9/FolVPS21 | 6.2 ± 0.1   | 3.0 ± 0.2    | 0.6 ± 0.1    | 9.6 ± 1.5           |

*SD was calculated from three repeated experiments and asterisks indicate statistically significant differences (p < 0.01), NA, not assay.

**Figure 1** FolVps9 is involved in vegetative hyphal growth and asexual development. (A) Colony morphology. Wild type, ΔFolvps9 mutant and the complemented strains were grown on CM and PDA media at 28°C for 5 days in the dark. (B) Conidiation. The indicated strains were cultured in CMC liquid medium for 5 days and photographed under differential interference contrast (DIC) microscopy. Bar = 10 µm. C, conidium. (C) Microconidia morphology. Wild type, ΔFolvps9 mutant and the complemented strains were cultured in CMC liquid medium for 5 days and observed by DIC microscopy. Bar = 5 µm.
a RFP-FolVPS21 expression vector and co-transformed with the GFP-FolVps9 construct into Fol. The resulting neomycin-resistant transformants were screened by detection of the GFP and RFP signals using fluorescence microscopy. We found that a large proportion of the GFP-FolVps9 and RFP-FolVps21 proteins were co-localized in conidia and hyphae (Yellow signal; Figures 3A,B). This result suggested that FolVps9 was at least partially localized in endosomes.

The co-localization of FolVps9 with an endosomal marker such as FolVps21 indicates that it may function during endocytosis. To determine whether FolVps9 is involved in endocytosis, we used FM4-64, a fluorescent dye that stains phospholipids and has been used to track endocytosis in fungi, to investigate endocytosis in the ΔFolvps9 mutant (Fischer-Parton et al., 2000; Qi et al., 2016). In the wild-type and complemented strains, the FM4-64 stain was gradually internalized and stained endosomes were visible 1 min after incubation (Figures 3C,D). In contrast, only a weak fluorescent signal was visible on the plasma membrane in the ΔFolvps9 mutant after 1 min incubation, and endosomal compartments were not stained until 5 min (Figures 3C,D). These results suggested that endocytosis was delayed in the ΔFolvps9 mutant.

The Vps9 Domain of FolVps9 Is Required for Its Normal Localization and Function

To investigate the contribution of each domain (Vps9 and Cue) of FolVps9 to subcellular localization and function, we generated GFP-FolVps9ΔVps9 and GFP-FolVps9ΔCue constructs lacking the Vps9 and Cue domain, respectively, and transformed each construct into the ΔFolvps9 mutant (Figure 4A). The GFP signal of FolVps9ΔCue was localized to punctate structures (presumptive endosomes), similar to GFP-FolVps9 (Figure 4B). However, the green fluorescence of FolVps9ΔVps9 was distributed throughout the cytoplasm (left) and excluded from vacuoles (right) (Figure 4B).

Next, we assayed possible functions of the Vps9 and Cue domains in fungal development and plant infection. In contrast to the wild type and ΔFolvps9/FolVPS9 strains, FolVps9ΔVps9 exhibited phenotypes similar to the ΔFolvps9 mutant, whereas

FIGURE 2 | Folvps9 is required for pathogenicity in Fol. (A) Pathogenicity assays. The susceptible tomato cultivar Moneymaker was inoculated with wild-type, ΔFolvps9 mutant and complemented strains using a standard root-dip method. Photographs were taken 21 days after infection. (B) Vascular discoloration of tomato plants. Tomato plants were inoculated with conidial suspensions and then cultivated for 14 days, after which the stems were split and photographed using a stereo microscope. Bar = 1 mm. (C) Outgrowth of fungi from tomato stems of plants inoculated with the indicated strains on PDA medium. Plates were incubated for 2 days.
FolVps9ΔCue did not possess obvious defects (Figures 4C,D and Table 1). These results suggest that the Vps9, but not the Cue, domain is required for the correct localization and physiological functions of FolVps9.

**FolVps9 Is Essential for Autophagy in Fol**

During the process of autophagy, the autophagosomes are transported to vacuoles for degradation via vesicle trafficking. The autophagy process can be analyzed by monitoring the vacuolar delivery and subsequent breakdown of GFP-Atg8 by vacuolar hydrolases (Liu et al., 2015). To test whether FolVPS9 plays a role in autophagy, we introduced the autophagy marker fusion gene GFP-FolAtg8 into wild type and the ΔFolvps9 mutant. We analyzed the strains for autophagy flux using epifluorescence microscopy and also examined the autophagy bodies of hyphal cells grown in rich medium (CM) or minimal medium without NaNO₃ (MM-N) by transmission electron microscopy.

When grown in CM rich medium for 16 h, no autophagy bodies or GFP-FolAtg8 fluorescence signals appear in punctuate structures, and the surrounding CMAC stained vacuoles are distributed throughout the cytoplasm of the wild-type and ΔFolvps9 mutant strains (Figures 5A–C). When hyphae were shifted to nitrogen-starvation conditions (MM-N) for 8 h, wild type exhibited reduced levels of full-length GFP-FolAtg8 and a more abundant free GFP band (Figure 5D). In contrast, compared to wild type, levels of full length GFP-FolAtg8 were relatively greater in the ΔFolvps9 mutant on MM-N medium (Figure 5D). Thus, the GFP-FolAtg8 assays demonstrated that FolVps9 is required for normal GFP-FolAtg8 proteolysis. Taken together, our results suggest that FolVps9 is important for autophagy in Fol.

**FolVps9 Interacts With the GDP-Bound Form of FolVps21**

In order to screen for potential regulators of endocytosis and autophagy in response to FolVps9, we used the FolVps9-3 X FLAG strain to identify co-immunoprecipitated (IP)
proteins by mass spectrometry (Table 2). Proteins that co-immunoprecipitated with FolVps9 include the Rab GTPase FolYpt1 and a number of vesicle trafficking proteins (Table 2). One of the FolVps9 interacting proteins, FolVps21, is highly similar to yeast Vps21p that functions in endocytosis and autophagy. Vps21 is a small Ras-related GTPase that can continuously cycle between the cytosol and plasma membrane by converting between its GDP-bound and GTP-bound form (Krick et al., 2008). Several vps21 mutations that lead to constitutive occupancy with GDP or GTP have been well studied in yeast and M. oryzae: for example, the S21L/S25N mutation (GDP-bound form) and Q66L/Q70L mutation (GTP-bound form) in yeast and M. oryzae, respectively (Lo et al., 2011; Zhu et al., 2018). Based on this precedent, we examined wild type FolVps21 and the FolVps21S27N (predicted GDP-bound form) and FolVps21Q72L (predicted GTP-bound form) mutations.

We first used the yeast two-hybrid (Y2H) assay to detect interactions between FolVps9 and the three states of FolVps21. The Y2H results showed that FolVps9 specifically interacted with FolVps21S27N, but not with FolVps21 or FolVps21Q72L (Figure 6A), indicating that FolVps9 might act as a FolVps21 GEF. We next explored the role of each domain (Vps9 and Cue) of FolVps9 in the interaction with FolVps21, by generating Y2H constructs expressing FolVps9ΔVps9 and FolVps9ΔCue and assaying their interaction with FolVps21S27N by Y2H. The results revealed that the Vps9 domain was required for the interaction of FolVps9 with FolVps21S27N (Figure 6A).

The Y2H interaction between FolVps9 and FolVps21S27N was corroborated using BiFC and co-IP assays in Fol (Figures 6B,C). Fluorescence was only observed in a Fol strain expressing both FolVps9-YFP and FolVps21S27N-YFP (Figure 6B). Taken together, these results indicate that FolVps9 interacts with FolVps21S27N, but not with wild type FolVps21 or the GTP-bound form, FolVps21Q72L.

**FolVps21 Is Important for Growth Polarity, Conidiation, and Full Virulence**

Because FolVps21 physically interacts with FolVps9, we further characterized the biological role of FolVPS21 using a targeted gene replacement strategy in Fol (Supplementary Figures S1A,C). The ΔFolvs21 gene deletion mutant displayed...
very similar phenotypes to the ΔFolvps9 mutant. The polarized growth rate was slower in the ΔFolvps21 mutant (Figure 7A and Table 1). Microscopic observations showed that the ability to produce conidia was reduced in the ΔFolvps21 mutant (Figure 7B). After culturing in CMC medium for 5 days, the ΔFolvps21 mutant produced $1.5 \times 10^8$ microconidia/ml compared to $3.1 \times 10^8$ microconidia/ml produced by the wild-type strain (Table 1). Furthermore, 78.3% of the conidia produced by the ΔFolvps21 mutant were smaller than those from the wild type (Table 1). In addition, the ΔFolvps21 mutant exhibited reduced macroconidia and virulence on tomato (Table 1 and Figures 7D,E, Supplementary Figure S2). The defects in development and pathogenicity were fully restored in the complemented strain (Figure 7 and Table 1). Taken together, these results suggest that FolVps9 and FolVps21 function together in mediating the same physiological process in Fol.

### Table 2

| Gene ID     | Description                                      | Homology in yeast | Unique peptides | MW (kDa) |
|------------|--------------------------------------------------|-------------------|-----------------|----------|
| FOXG_02298 | Small COP-II coat GTPase                          | Sar1              | 2               | 21.5     |
| FOXG_03079 | Myosin heavy chain                                | Myo1              | 4               | 275.5    |
| FOXG_12799 | Syntaxin 7                                        | Vam3              | 2               | 29.7     |
| FOXG_02392 | Rabs family GTPase                                | Vps21             | 3               | 24.7     |
| FOXG_03741 | Golgi transport complex subunit                   | Cog6              | 2               | 48.5     |
| FOXG_12808 | GTP-binding protein ypt1                          | Ypt1              | 3               | 22.4     |
| FOXG_07785 | Hypothetical protein                              | Vps41             | 4               | 148.9    |
| FOXG_04244 | Syntaxin 16                                       | Tlg2              | 3               | 38.6     |
| FOXG_02138 | Hypothetical protein                              | Muk1              | 2               | 76.8     |

**Constitutively Activated FolVps21 Rescues the Defects of the ΔFolvps9 and ΔFolvps21 Mutants**

Numerous studies in yeast suggest that Vps9p functions as a highly efficient GEF and positive regulator for Vps21p (Esters et al., 2001; Bean et al., 2015; Zhu et al., 2018). Our results demonstrating that FolVps9 only interacts with the predicted GDP-bound form of FolVps21 and that mutants lacking either gene possess the same phenotypes supports a similar scenario in Fol. To further investigate whether FolVps9 might function as a positive regulator/GEF for FolVps21 in
**FIGURE 6** | FolVps9 interacts with the predicted GDP-bound form of FolVps21. (A) Yeast two-hybrid assays. Yeast cells expressing the prey and bait constructs were assayed for growth on SD-Leu-Trp and SD-Leu-Trp-His-Ade plates. +, positive control; -, negative control. (B) Bimolecular fluorescence complementation (BiFC) analysis. Mycelia of transformants expressing the FolVps9-YFP/FolVps21S27N-YFPN constructs were observed by DIC or fluorescence microscope. Bar = 10 µm. (C) Co-immunoprecipitation assays. Total proteins were extracted from transformants co-expressing FolVps9-3xFLAG and GFP-FolVps21S27N constructs, and protein eluted from the anti-GFP M2 beads (elution). Immunoblots were incubated with monoclonal anti-FLAG or anti-GFP antibody, as indicated.

Fol, we generated constitutively active FolVPS21Q72L (GTP-bound form) and dominant negative FolVPS21S27N (GDP-bound form) constructs under the control of the RP27 promoter and transformed them into protoplasts of the ΔFolvps9 mutant. The resulting ΔFolvps9/FolVPS21Q72L transformants were examined expression of FolVps21 by qRT-PCR analysis (Supplementary Figure S3). Phenotypic analysis revealed that the ΔFolvps9/FolVPS21Q72L transformant displayed normal vegetative growth, conidiation, conidial morphology, subcellular localization and virulence compared to wild type (Figure 8 and Table 1). In contrast, the ΔFolvps9/FolVPS21S27N strain resembled the ΔFolvps9 mutant. These results support a positive role for FolVps9 in regulating the biological function of FolVps21, consistent with a function as a GEF for FolVps21.

We also transformed FolVPS21S27N and FolVPS21Q72L constructs into protoplasts of the ΔFolvps21 mutant. The resulting ΔFolvps21/FolVPS21Q72L transformants were examined by qRT-PCR analysis (Supplementary Figure S3). Phenotypic analysis gave results very similar to those observed for the FolVPS21S27N and FolVPS21Q72L constructs transformed into the ΔFolvps9 mutant. The data revealed that growth, conidiation and pathogenicity defects associated with the ΔFolvps9 mutation were fully rescued by FolVPS21Q72L (Figure 9 and Table 1). However, the FolVPS21S27N allele was not able to rescue the phenotype defects of the ΔFolvps21 mutant (Figure 9 and Table 1). In addition, the localization pattern of RFP-tagged FolVps21Q72L in the ΔFolvps21/RFP-FolVPS21Q72L strain was similar to that observed in the ΔFolvps21/RFP-FolVps21 strain (Figure 9). The localization pattern of RFP-tagged FolVps21Q72L in the ΔFolvps21/RFP-FolVPS21S27N strain was altered compared to that of the wild type protein observed in the ΔFolvps21/RFP-FolVps21 strain (Figure 9). Taken together, these results showed that constitutively activated FolVps21 can rescue the growth,
Li et al. Endocytosis and Autophagy in Fusarium oxysporum

FIGURE 7 | FolVps21 is important for growth polarity, asexual development and full virulence. (A) Colony growth and morphology assays were performed as described in Figure 1. Bar = 10 µm. C, conidium. (B) Conidiophore development assays were performed as described in Figure 1. (C) Conidial morphology. Conidia from wild-type, ΔFolvps21 mutant and the complemented strains were observed by DIC microscopy. Bar = 5 µm. (D) Moneymaker tomato plants were infected with the indicated strains as described in Figure 2 and photographs were taken at 21 dpi. (E) Vascular discoloration. Tomato plants were inoculated with conidial suspensions for 14 days, after which the stems were split and photographed using a stereomicroscope. Bar = 1 mm.

Constitutively Activated FolVps21 Corrects the Endocytosis and Autophagy Defects of the ΔFolvps9 Mutant

As shown above, phenotypic data indicates that constitutively FolVps21 can rescue ΔFolvps9 defects. In order to test the effects of constitutively activated FolVps21 on endocytosis impairments in the ΔFolvps9 mutant, we performed FM4-64 dye uptake experiments. The defects in FM4-64 uptake by the ΔFolvps9 mutant were suppressed by expression of constitutively activated FolVps21, but not by constitutively inactive FolVps21 (Figure 10). The ΔFolvps9/FolVPS21Q72L (85.8%) strain exhibited relatively faster FM4-64 uptake (1 min) than the ΔFolvps9 mutant (5 min), while FM4-64 uptake in the ΔFolvps9/FolVPS21S27N (84.8%) strain was similar to the ΔFolvps9 mutant (5 min) (Figure 10). These results suggest that constitutively activated FolVps21 suppresses the endocytosis defects of the ΔFolvps9 mutant.

We also utilized transmission electron microscopy to observe autophagy bodies of hyphal cells in our strains. When grown in CM medium, no autophagy bodies were observed in the hyphae of any strain (Figure 11A). However, when hyphae were shifted to MM-N medium, numerous autophagy bodies accumulated in the vacuoles of wild-type and ΔFolvps9/FolVPS21Q72L strains (Figure 11A).

We next analyzed the GFP-FolAtg8 fluorescence signals under two culture conditions in our strains. In CM medium, the GFP-FolAtg8 fluorescence signal existed in punctuate structures and the surrounding CMAC-stained vacuoles were distributed in the cytoplasm of all strains (Figure 11B). After culture in MM-N medium, GFP fluorescence signals accumulated in the vacuoles.

developmental and pathogenicity defects of the ΔFolvps9 and ΔFolvps21 mutants in Fol.
of wild-type and ΔFolvps9/FolVPS21Q72L strains (Figure 11B). However, little GFP fluorescence and few autophagy bodies were observed in the vacuoles of the ΔFolvps9 mutant and ΔFolvps9/FolVPS21S27N strains (Figures 11B,D). To further confirm these observations, we also performed a GFP-FolAtg8 proteolysis assay. Under normal conditions, a clear full length GFP-FolAtg8 fusion protein band (40 kDa) could be detected in all strains (Figure 11C). When hyphae were shifted to nitrogen-starvation conditions, the wild-type and ΔFolvps9/FolVPS21Q72L strains displayed relatively weak full-length GFP-FolAtg8 and an abundant GFP band (Figure 11C). Compared to wild type, levels of full length GFP-FolAtg8 did not significantly decrease in the ΔFolvps9 and ΔFolvps9/FolVPS21S27N strains (Figure 11C). These results support the hypothesis that constitutively activated FolVps21 can rescue autophagy defects of the ΔFolvps9 mutant.

**DISCUSSION**

Guanine nucleotide exchange factors (GEFs) facilitate release of GDP and the binding of GTP for subsequent activation of Rab GTPases (Ishida et al., 2016). The Vps9 GEF protein and the Vps21 GTPase have been reported to play important roles in modulating development in organisms from yeast to humans (Esters et al., 2001; Paulsel et al., 2013). However, the autophagy and endocytosis functions of Vps9 protein have been investigated in a limited number of studies in filamentous fungi, and with only one plant fungal pathogen (M. oryzae). In this study, we have identified and characterized the Vps9 homolog FolVps9 in the tomato pathogen Fol and showed that FolVps9 was required for fungal growth, development, endocytosis, autophagy and plant pathogenicity. Based on genetic evidence and cytological examination, we further obtained evidence supporting a role for FolVps9 as a GEF for FolVps21, by revealing a physical association between the two proteins and showing that constitutively activated FolVps21 was able to rescue the defects of the ΔFolvps9 mutant. In addition, we demonstrated that FolVps21 is also important for fungal growth, development and plant infection, and that the FolVps9-FolVps21Q72L interaction plays an important role in endocytosis and autophagy of Fol (Figure 12).

In this study, we determined that the ΔFolvps9 mutant had striking defects in polarity growth, conidiogenesis and...
FIGURE 9 | Constitutively activated FolVps21 (Q72L) rescues the defects of the ΔFolVps21 mutant strain. (A) Colony and conidial morphology assays were performed as described in Figure 1. Bar = 10 µm. (B) Subcellular localization of the indicated different states of the FolVps21 protein. Bar = 10 µm. (C) Pathogenicity of mutants in tomato. Moneymaker tomato seedlings were inoculated with conidial suspensions of the indicated strains as described in Figure 2 and photographs were taken at 21 dpi.

FIGURE 10 | Constitutively activated FolVps21 (Q72L) corrects the endocytosis defects of the ΔFolVps9 mutant strain. Strains were grown in liquid CM medium for 48 h, then stained with FM4-64 and examined using a confocal fluorescence microscope (Zeiss LSM710, 63x oil objective) as described in Figure 3. The camera exposure is indicated in milliseconds (800 ms). The numbers indicate the percentage of hyphae exhibiting the endocytosis pattern seen in the figure (n = 100). Bar = 10 µm.

pathogenicity. Consistent with these results, in *F. graminearum*, Vps39, a HOPS complex subunit, localizes to early endosomes (EEs) and plays key roles in endocytosis and morphology (Li B. et al., 2017). Other studies have shown that endocytosis was important for the pathogenic development of different plant fungal pathogens (Li L. et al., 2017; Li X. et al., 2017;
Constitutively activated FolVps21 (Q72L) rescues the autophagy defects of the δFolvps9 mutant strain. (A) Examination of organelles and autophagic bodies by transmission electron microscopy as described in Figure 5. Bar = 1 µm. C, cytoplasmic; V, vacuole. Asterisks indicate autophagosomes. (B) Localization of GFP-FolAtg8 and staining with 7-amino-4-chloromethylcoumarin (CMAC). The indicated Fol strains were assayed as described in Figure 5. Bar = 10 µm. (C) GFP-FolAtg8 proteolysis assays were performed as described in Figure 5. (D) Statistical analysis of 100 cells from the indicated strains (B) for GFP signal in the vacuole. Error bars represent ± SD and asterisks indicate statistically significant differences (p < 0.01).

Li et al., 2019b). Through systems-level analysis of the functions of endocytic proteins, such as the early endocytic protein MoEnd3 (Li X. et al., 2017), the SNARE protein FolVam7 (Li et al., 2019b), and the capping protein MoCap (Li L. et al., 2017), it was revealed that endocytosis was important for development and virulence in different plant fungal pathogens.

Autophagy is a fundamental function in eukaryotes and is well conserved from mammals to yeasts. Nevertheless, there is a little evidence to demonstrate any relationship between endocytosis and autophagy in filamentous fungi. Here, we found that FolVps9 co-localized with FolVps21 and that the δFolvps9 mutant exhibited defects in endocytosis. In addition, the FolVps9 mutant has defects in the trafficking and degradation of autophagosomes. Several Rab proteins were well-known mediators of vesicle trafficking and involved in regulation of autophagy (Liu et al., 2015; Zheng et al., 2018b). For example, Rab5 plays roles in vesicle trafficking, cargo recycling and maturation of early endosomes and is involved in membrane
fusion events. Similar to its endosome function, Rab5 can recruit the autophagic BECN1-PI3K3C3 complex and production PI3P on early stage of autophagosomal formation (Ravikumar et al., 2008). Rab1 regulates vesicle trafficking and functions during early autophagy (Zoppino et al., 2010).

In Fol, we did not able to isolate the ΔFolvps9 and ΔFolvps21 double mutant and it was thus difficult to directly demonstrate the relationship between autophagy and endocytosis. However, based on our functional analysis of FolVps9 and previous work, we speculate that there is crosstalk between autophagy and endocytosis. Here, we used mass spectrometry to identify several candidate vesicle trafficking and Rab GTPase proteins that interact with FolVps9, including FolYpt1, FolVps21, FolSar1, FolVam3, and FolVps41. Furthermore, our protein-protein interaction results confirmed that FolVps9 interacts with the GDP-bound form of FolVps21. Taken together, these results suggest that FolVps9 interacts with these candidate proteins to participate in autophagy and endocytosis in Fol.

The reduced virulence of the ΔFolvps9 mutant results from multiple defects. First, the vegetative growth of ΔFolvps9 mutant was significantly reduced relative to the wild type strain. Consistent with these results, the decreased growth of the ΔFolvps9 mutant led to attenuated virulence in the plant pathogenic fungus M. oryzae (Zhu et al., 2018). Second, the ΔFolvps9 mutant showed delayed endocytosis, which was regarded as one of fungal responses to plant infection (Li X. et al., 2017; Li B. et al., 2017). We obtained evidence that the impaired virulence of the ΔFolvps9 mutant was due to a defect in endocytosis, in that the impairment in virulence was suppressed when endocytosis was rescued in the ΔFolvps9 mutant. Third, we showed that perturbing the autophagy pathway causes attenuated virulence in the ΔFolvps9 mutant, as the autophagy pathway is well conserved and important during the invasion of host cells (Kershaw and Talbot, 2009; Lv et al., 2017). Following the constitutive activation of FolVps21 in the ΔFolvps9 mutant, defects in autophagy and virulence were suppressed, indicating that autophagy was important to infection for Fol.

Insights from a previous study indicate that the Vps9 domain is important for regulating protein-protein interactions in bacteria (Bair et al., 2008). In this study, we demonstrated that the Vps9 domain was required for the interaction between FolVps9 and FolVps21S27N. This result was consistent with what has been observed in M. oryzae and F. graminearum (Li et al., 2015; Zheng et al., 2018a; Zhu et al., 2018). More importantly, we found that while expression of constitutively active FolVps21 in the FolVPS9 deletion mutant could restore normal development and pathogenesis, the dominant negative form of FolVps21 could not rescue the defects associated with the ΔFolvps9 mutant. Taken together, our results indicate that FolVps9 likely functions as a GEF for FolVps21 to convert FolVps21-GDP to the GTP-bound form to promote vegetative growth, development and infection in Fol.

In summary, we identified and characterized FolVPS9, a gene encoding a putative vacuolar sorting protein in Fol, which...
plays crucial role in hyphal growth, conidiation, endocytosis, autophagy and pathogenesis. We also provide evidence that FolVps9 might function as a GEF for FolVps21. Our studies provide preliminary evidence for crosstalk between autophagy and endocytosis in Fol. Further research efforts will be focused on investigation of the precise molecular function of FolVps9 and its associated proteins and how they regulate endocytosis and autophagy. Results from our studies also provide targets for developing sustainable control strategies for fungal plant pathogens.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/Supplementary Material.

**AUTHOR CONTRIBUTIONS**

S-QO and BL conceived and designed the experiments and wrote the manuscript. BL, H-YM, and Z-YZ performed the experiments. BL, X-JC, and S-QO analyzed the experiment data. All authors have read and approved the final manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.02658/full#supplementary-material

**FIGURE S1** | Generation of the FolVPS9 and FolVPS21 deletion mutants. (A) Schematic diagram of the FolVPS9 and FolVPS21 genes, and split-marker gene replacement constructs. HPH-N and HPH-C are the N and C-terminal halves of the cassette that confers resistance to hygromycin B. (B,C) Southern blot analysis. Genomic DNA from wild type and presumptive deletion mutants was digested with EcoRI and HindIII subjected to Southern blot analysis using gene-specific and HPH probes, respectively, WT, Wild type.

**FIGURE S2** | qPCR to determine relative levels of Fol in stem sections of inoculated tomato plants. Genomic DNA was isolated from tomato stems of the Moneymaker cultivar infected with various Fol strains as described in section Materials and Methods. Quantitative PCRs were performed to evaluate the fungal biomass using primers that amplify the intergenic spacer region of the ribosomal 28S subunit. The mean values of three determinations with standard deviations are shown. Asterisks indicate statistically significant difference relative to wild type Fol (p < 0.01).

**FIGURE S3** | Expression of FolVPS21 in the indicated strains. The expression of FolVPS21 was measured by quantitative real-time PCR with CDN4 isolated from hyphae of the indicated strains. Levels were first normalized to actin. The relative abundance of FolVPS21 transcripts was then normalized to that in wild-type vegetative hyphae (Relative transcript level = 1). Error bars represent ± SD and asterisks indicate statistically significant differences (p < 0.01).

**TABLE S1** | Wild-type and mutant strains of fungi used in this study.

**TABLE S2** | Primers used in this study.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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