Dephosphorylation of the Core Septin, AspB, in a Protein Phosphatase 2A-Dependent Manner Impacts Its Localization and Function in the Fungal Pathogen Aspergillus fumigatus

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Septins are a conserved family of GTPases that form hetero–oligomeric complexes and perform diverse functions in higher eukaryotes, excluding plants. Our previous studies in the human fungal pathogen Aspergillus fumigatus revealed that the core septin, AspB, a CDC3 ortholog, is required for septation, conidiation, and conidial cell wall organization. Although AspB is important for these cellular functions, nothing is known about the role of kinases or phosphatases in the posttranslational regulation and localization of septins in A. fumigatus. In this study, we assessed the function of the Gin4 and Cla4 kinases and the PP2A regulatory subunit ParA, in the regulation of AspB using genetic and phosphoproteomic approaches. Gene deletion analyses revealed that Cla4 and ParA are indispensable for hyphal extension, and Gin4, Cla4, and ParA are each required for conidiation and normal septation. While deletion of gin4 resulted in larger interseptal distances and hypervirulence, a phenotype mimicking aspB deletion, deletion of cla4 and parA caused hyperseptation without impacting virulence, indicating divergent roles in regulating septation. Phosphoproteomic analyses revealed that AspB is phosphorylated at five residues in the GTPase domain (S134, S137, S247, T297, and T301) and two residues at its C-terminus (S416 and S461) in the wild-type, ∆gin4 and ∆cla4 strains. However, concomitant with the differential localization pattern of AspB and hyperseptation in the ∆parA strain, AspB remained phosphorylated at two additional residues, T68 in the N-terminal polybasic region and S447 in the coiled-coil domain. Generation of nonphosphorylatable and phosphomimetic strains surrounding each differentially phosphorylated residue revealed that only AspB<sub>mt</sub>-T68E showed increased interseptal distances, suggesting that dephosphorylation of T68 is important for proper septation. This study highlights the importance of septin phosphorylation/dephosphorylation in the regulation of A. fumigatus hyphal septation.

Keywords: Aspergillus fumigatus, septin, phosphorylation, kinase, phosphatase
INTRODUCTION

Septins, a conserved family of GTPases, are involved in a variety of critical cellular functions, ranging from cell division to cell wall maintenance (Momany et al., 2001; Alvarez-Tabares and Perez-Martin, 2010; Kozubowski and Heitman, 2010; Lindsey et al., 2010; Hernandez-Rodriguez et al., 2012, 2014). The number of septin-encoding genes varies amongst all organisms. In the pathogenic fungus Candida albicans there are seven septin genes, all of which have orthologs in the model yeast Saccharomyces cerevisiae, while Aspergillus fumigatus has only five septin genes (Warenka and Konopka, 2002; Pan et al., 2007; Juvvadi et al., 2011b). A. fumigatus aspA, aspB, aspC, and aspD are orthologous to S. cerevisiae CDC11, CDC3, CDC12, and CDC10, respectively; while aspE is absent in S. cerevisiae (Pan et al., 2007; Lindsey et al., 2010; Juvvadi et al., 2011b). Deletion analyses in A. fumigatus revealed that none of the septins are required for growth under basal conditions, yet septins AspA, AspB, AspC, and AspE play a key role in regular septation and AspA, AspB, AspC, and AspD regulate conidiation (Vargas-Muniz et al., 2015). Deletion of aspB resulted in hypervirulence in an invertebrate model of invasive aspergillosis, as well as increased susceptibility and AspB mislocalization following exposure to anti-cell wall agents.

Septins function through formation of heteropolymeric complexes; however, regulation of the formation of these complexes is unclear (Gladfelter, 2010). Multiple lines of evidence suggest that posttranslational modification, such as phosphorylation, regulates the formation, and stability of septin complexes. For example, posttranslational modification of septins in S. cerevisiae is key in controlling the assembly of septins into higher-order structures (Tang and Reed, 2002; Garcia et al., 2011). Septin Cdc3, Cdc10, Cdc11, Cdc12, and Shs1 are also phosphorylated in the filamentous hemiascomycete Ashbya gossypii (Mesoroll et al., 2013). Phosphomimetic mutation of A. gossypii septin Cdc12 and Shs1 phosphorylation sites resulted in lethality, supporting the notion of septin phosphorylation as a key regulator of proper septin function (Mesoroll et al., 2013).

Kinases Gin4, and Cla4 have been shown to phosphorylate septins in hemiascomycetes (Longtine et al., 1998; Kadota et al., 2004; Versele and Thorner, 2004; Wightman et al., 2004; Sinha et al., 2007; DeMay et al., 2009; Li et al., 2012). In S. cerevisiae, Gin4 co-purifies with and is regulated by the septins (Barral et al., 1998; Mortensen et al., 2002). Gin4 requires this interaction with the septin complex in order to be activated by hyperphosphorylation and subsequently phosphorylates the non-core septin SHS1 (Barral et al., 1998; Mortensen et al., 2002). This co-dependence between Gin4 and septins has also been demonstrated in C. albicans, where deletion of gin4 eliminates septin ring formation (Wightman et al., 2004). Another important regulator of septins in S. cerevisiae is the PAK kinase Cla4. In the S. cerevisiaeΔcla4 strain, septins form a band at the tip of the bud instead of localizing at the bud neck (Cvrekova et al., 1995). Furthermore, Cla4 is capable of interacting in vitro with and phosphorylating septins Cdc3 and Cdc10. Cla4 phosphorylates Cdc10 on two serine residues (S256 and S312), and alanine substitution of serine 256 resulted in an elongated bud at 37°C, indicating that this phosphorylation site is required for proper Cdc10 function (Versele and Thorner, 2004). Although the essential roles of Gin4 and Cla4 in septin phosphorylation have been described in hemiascomycetes, their role as potential septin regulators has not been explored in filamentous ascomycetes. Recent work in the model filamentous fungi Neurospora crassa and A. nidulans has shown that Cla4 is involved in hyphal extension, polarity, and asexual and sexual reproduction (Park et al., 2011; De Souza et al., 2013). The filamentous plant pathogen Magnaporthe oryzae cla4 ortholog (CHM1) is also involved in hyphal extension, polarized growth, new growth foci limitation, conidiation and pathogenesis (Li et al., 2004). In the basidiomycete Ustilago maydis, cla4 is involved in budding, cytokinesis, cell wall assembly, mating and pathogenesis (Leveleki et al., 2004).

Although some septin kinases have been explored, less is known of septin phosphatases. In S. cerevisiae, Rts1, a protein phosphatase 2A (PP2A) subunit, regulates septin dephosphorylation during telophase, and this dephosphorylation contributes to cytokinesis (Dobbelare et al., 2003). In A. nidulans, deletion of parA (RTSI ortholog) resulted in hyperseptation and reduction in conidiation; however, the possible septin regulatory role was not defined (Zhong et al., 2014).

In this study, we built on our previous explorations of septins in the pathogen A. fumigatus and defined for the first time the phosphorylation status of the core septin AspB and the role of Gin4, Cla4, and ParA as AspB regulators in A. fumigatus growth and development. We found that ParA and Cla4 are indispensable for full hyphal extension, whereas Gin4, Cla4, and ParA are important for conidiation and required for proper localization of AspB. While deletion of gin4 resulted in increased intersepal distances, deletion of cla4 or parA resulted in hyperseptation in more basal compartments. Similar to the ΔaspB strain, the Δgin4 strain is hypervirulent in the Galleria mellonella invertebrate and murine models of invasive aspergillosis. Phosphoproteomic analyses revealed that AspB is phosphorylated in seven residues, five of which are located in the GTPase domain. Interestingly, although the deletion of kinases gin4 or cla4 did not affect the phosphorylation status of AspB, the deletion of the phosphatase subunit parA resulted in the phosphorylation of two additional sites, including one residue (S447) in the coiled-coil domain. Mutation of these residues to alanine (blocking phosphorylation) or glutamic acid (phosphomimetic) mislocalized AspB and caused an increase in intersepal distances in the AspBΔT68E strain. Taken together, these important findings suggest that ParA is a potential direct regulator of AspB through dephosphorylation, while Cla4 and Gin4 could regulate AspB through a yet unknown indirect mechanism.

MATERIALS AND METHODS

Strains, Media, and Culture Conditions

The A. fumigatus akuBΔKU80 pyrGΔ uracil/uridine auxotroph was used for deletion analyses and the akuBΔKU80 uracil/uridine prototroph was used as the wild-type strain.
Aniline blue staining was used for detection of cell wall β-glucan. 10^4 conidia of each strain were cultured on coverslips immersed in 10 ml of GMM + UU broth and incubated for 20 h at 37°C, as previously described (Juvvadi et al., 2011a). The coverslips were rinsed with GMM + UU broth, inverted over 500 µl of aniline blue stain, and incubated for 5 min at 25°C. Coverslips were rinsed once more briefly with GMM + UU broth and observed by fluorescence microscopy.

**Galleria mellonella and Murine Invasive Aspergillosis Virulence Models**

Virulence of the deletion strains was assessed in an iterative fashion using an invertebrate and then a murine model of invasive aspergillosis. For the initial invertebrate model, 20 larvae of the wax moth *G. mellonella* were infected with each of the *A. fumigatus* deletion strains or the wild-type strain, delivering 5 µl of a 1 × 10^6 conidia/ml of suspension with a total inoculum size of 2 × 10^5 conidia. Infected larvae were incubated at 37°C and survival scored daily for 5 days (Steinbach et al., 2006). For the murine model, male mice (CD1, Charles River Laboratory, Raleigh, NC, USA) were immunosuppressed with cyclophosphamide (175 mg/kg, intraperitoneally, days −2 and +3) and triamcinolone acetonide (40 mg/kg, subcutaneously, days −1 and +6). A total inoculum of 4 × 10^6 conidia was delivered intranasally using 40 µl of 10^8 conidia of the Δgin4, Δcla4, ΔparA, and ΔkuB strains (Vargas-Muniz et al., 2015). Survival for both virulence models was plotted on a Kaplan–Meier curve and analyzed using log rank pair-wise comparison. Animal studies were carried out in accordance with all of the guidelines of the Duke University Medical Center Institutional Animal Care and Use Committee (IACUC) and in compliance with the United States Animal Welfare Act (Public Law 98-198). Duke University Medical Center IACUC approved all of the vertebrate studies. The studies were

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**Construction of Septin Deletion Strains**

Deletion of *cla4* was performed by replacing the 2.6 kb *cla4* gene (Afu5g05900) with the 2.4 kb *pyrG* gene from *A. parasiticus*. Approximately 1 kb of upstream and downstream flanking regions of *cla4* were PCR-amplified from AF293 genomic DNA (Supplementary Table S1). *pyrG* was amplified from the pJW24 plasmid. The *cla4* deletion construct was generated by fusion PCR and transformed into the Δ*kuB* strain, as previously described (Steinbach et al., 2006). Deletion of *gin4* was performed by similarly replacing the 4.0 kb *gin4* gene (Afu6g02300) with the 2.4 kb *pyrG* gene from *A. parasiticus*. Deletion of *parA* was attained by replacing the 2.5 kb *parA* gene (Afu5g02560) with the 3.0 kb *A. parasiticus* *pyrG* cassette, and the resulting plasmid was digested with NsiI and SalI and transformed into Δ*kuB* strains. Each deletion strain was then transformed with the *aspB*–*egfp* construct as previously described (Vargas-Muniz et al., 2015). All gene deletions were confirmed by both PCR (data not shown) and Southern analyses (Supplementary Figure S1).

**Radial Growth and Conidial Quantification**

Conidia (10^4) from each strain were inoculated on GMM agar, incubated at 37°C, and radial growth measured every 24 h for 5 days. For conidial quantification, 10^4 conidia from each strain were inoculated on GMM agar, incubated for 5 days at 37°C, and conidia harvested in 10 ml of 0.05% Tween-80 and quantified using a hemocytometer as previously described (Lamoth et al., 2012). All assays were performed in triplicate. Student’s t-tests were performed using Graph Pad Prism (San Diego, CA, USA).  

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**TABLE 1 | Strains used in the present study.**

| Strain | Parent strain | Genotype | Reference on source |
|--------|---------------|----------|---------------------|
| ΔkuB* | CEA17 pyrG+ | pyrG | da Silva Ferreira et al., 2006 |
| ΔkuB* | CEA17 | Wild-type | da Silva Ferreira et al., 2006 |
| aspB–egfp | ΔkuB* | ΔaspB:aspBpromo-aspB–egfp-hph | Vargas-Muniz et al., 2015 |
| Δgin4 | ΔkuB* | Δgin4::pyrG | This study |
| Δcla4 | ΔkuB* | Δcla4::pyrG | This study |
| ΔparA | ΔkuB* | ΔparA::pyrG | This study |
| Δgin4 ΔaspB | Δgin4 | Δgin4::pyrG ΔaspB::ble | This study |
| Δgin4 aspB–egfp | Δgin4 | Δgin4::pyrG ΔaspB:aspBpromo-aspB–egfp-hph | This study |
| Δcla4 aspB–egfp | Δcla4 | Δcla4::pyrG ΔaspB:aspBpromo-aspB–egfp-hph | This study |
| ΔparA aspB–egfp | ΔparA | ΔparA::pyrG ΔaspB:aspBpromo-aspB–egfp-hph | This study |
| aspB*::Δ447A | ΔkuB* | ΔaspB:aspBpromo-aspB::Δ447A-egfp-hph | This study |
| aspB*::Δ447E | ΔkuB* | ΔaspB:aspBpromo-aspB::Δ447E-egfp-hph | This study |
| aspB*::T68A | ΔkuB* | ΔaspB:aspBpromo-aspB::T68A-egfp-hph | This study |
| aspB*::T68E | ΔkuB* | ΔaspB:aspBpromo-aspB::T68E-egfp-hph | This study |

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(da Silva Ferreira et al., 2006; Table 1). Cultures were grown on glucose minimal media (GMM) at 37°C, except where otherwise specified. *Escherichia coli* DH5α competent cells were used for cloning.
conducted in the Division of Laboratory Animal Resources (DLAR) facilities that are accredited by the Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC).

**Histopathology Analyses**
To characterize *in vivo* disease histopathology, three additional mice per each single deletion strain were infected. Mice were euthanized on day +3 after inoculation and lungs were harvested. Lungs sections were stained with Gomori’s methenamine silver stain to stain fungal hyphae and hematoxylin and eosin stain to examine inflammation (Steinbach et al., 2006).

**Fluorescence Microscopy**
Conidia (10^6) of each *aspB-egfp* expressing strain were cultured on coverslips immersed in 10 ml of GMM broth and incubated for 20 h at 37°C, as previously described (Vargas-Muniz et al., 2015). Localization patterns were visualized using an Axioskop 2 plus microscope (Zeiss) equipped with AxioVision 4.6 imaging software.

**Protein Extraction, AspB-EGFP Fusion Protein Purification and LC–MS/MS Analysis**
Each *A. fumigatus* strain expressing the *aspB-egfp* fusion construct under the control of the *aspB* native promoter was grown in GMM broth for 24 h at 37°C. Biological replicates were prepared for each strain. Total cell lysate was obtained by homogenizing mycelia (600–650 mg wet weight) as previously described (Juvvadi et al., 2013). Total protein in the crude extract was quantified by the Bradford method and normalized to contain 10 mg of protein in the sample before purification using GFP-Trap® affinity purification (Chromotek). GFP-Trap® resin was equilibrated according to the manufacturer’s instructions and resuspended in 100 µl ice-cold dilution buffer (10 mM Tris–HCl pH 7.5, 150 mM NaCl, 0.5 mM EDTA, 1 mM PMSF, 1:100 protease inhibitor cocktail). GFP-Trap® resin suspension was then added to the crude lysate containing the 10 mg of protein and incubated for 2 h at 4°C with gentle agitation. The suspension was centrifuged at 2000 rpm for 10 min at 4°C and the GFP-Trap® pellet was washed once with 500 µl of iced-cold dilution buffer and twice with 500 µl of wash buffer (10 mM Tris–HCl pH 7.5, 350 mM NaCl, 0.5 mM EDTA, 1 mM PMSF, 1:100 protease inhibitor cocktail). The resin was finally washed 3 times with 200 µl of 50 mM ammonium bicarbonate and resuspended in 30 µl of 50 mM ammonium bicarbonate.

Protein bound GFP-Trap® resins suspended in 30 µl 50 mM ammonium bicarbonate, pH 8.0, were supplement with 0.1% Rapigest SF surfactant (Waters Corp). Samples were reduced with 5 mM dithiothreitol for 30 min at 70°C and free sulfhydryls were alkylated with 10 mM iodoacetamide for 45 min at room temperature. Proteolytic digestion was accomplished by the addition of 500 ng sequencing grade trypsin (Promega) directly to the resin with incubation at 37°C for 18 h. Supernatants were collected following a 2 min centrifugation at 1,000 rpm, acidified to pH 2.5 with TFA and incubated at 60°C for 1 h to hydrolyze remaining Rapigest surfactant. Insoluble hydrolyzed surfactant was cleared by centrifugation at 15,000 rpm for 5 min. The sample was lyophilized to dryness and phosphopeptides were enriched using GL Biosciences p10 TiO2 derivatized tips according to manufacturer protocols. Extracted peptides were lyophilized to dryness and resuspended in 12 µL of 0.2% formic acid/2% acetonitrile.

Each sample was subjected to chromatographic separation on a Waters NanoAcquity UPLC equipped with a 1.7 µm BEH130 C18 75 µm I.D. × 250 mm reversed-phase column. The mobile phase consisted of (A) 0.1% formic acid in water and (B) 0.1% formic acid in acetonitrile. Following a 4 µL injection, peptides were trapped for 3 min on a 5 µm Symmetry C18 180 µm I.D. X 20 mm column at 5 µL/min in 99.9% A. The analytical column was then switched in-line and a linear elution gradient of 5% B to 40% B was performed over 30 min at 400 nL/min. The analytical column was connected to a fused silica PicoTip emitter (New Objective, Cambridge, MA, USA) with a 10 µm tip orifice and coupled to a QExactive Plus mass spectrometer through an electrospray interface operating in a data-dependent mode of acquisition. The instrument was set to acquire a precursor MS scan from m/z 375–1675 with MS/MS spectra acquired for the ten most abundant precursor ions. For all experiments, HCD (higher energy collisional dissociation) energy settings were 27 v and a 120 s dynamic exclusion was employed for previously fragmented precursor ions.

Raw LC–MS/MS data files were processed in Proteome Discoverer (Thermo Scientific) and then submitted to independent Mascot searches (Matrix Science) against a RefSeq *Aspergillus* database containing both forward and reverse entries of each protein. Search tolerances were 5 ppm for precursor ions and 0.02 Da for product ions using trypsin specificity with up to two missed cleavages. Carbamidomethylation (+57.0214 Da on C) was set as a fixed modification, whereas oxidation (+15.9949 Da on M), deamidation (+0.98 Da on NQ), and phosphorylation (+79.99 Da on STY) was considered dynamic mass modifications. All searched spectra were imported into Scaffold (v4.3, Proteome Software) and scoring thresholds were set to achieve a peptide false discovery rate of 1% using the PeptideProphet algorithm. Peak area calculations from extracted ion chromatograms were generated within Skyline v3.5 (MacCoss Lab, University of Washington) following manual peak integration based on identification of retention time and accurate mass.

**RESULTS**
**Cla4 and ParA Are Required for Hyphal Extension**
Our previous studies showed that *A. fumigatus* septins play a pleiotropic role in septation, conidiation, and response to anti-cell wall agents (Vargas-Muniz et al., 2015). In order to gain a further understanding into the regulation and function of AspB in septation, we characterized the possible regulators of AspB using a candidate approach. Based on the literature, we selected the non-essential kinases Gin4 and Cla4, as well as the protein...
Deletion of the A. fumigatus core septin AspB resulted in reduction of conidia production and delayed septation (Vargas-Muniz et al., 2015). To further understand the possible role of Cla4, Gin4, and ParA as septin regulators, we measured conidial production in the respective deletion backgrounds (Figure 1B). The Δcla4, Δgin4, and ΔparA strains showed a 5.6-, 4.4-, and 20.7-fold, reduction in conidiation, respectively, compared to the akuB KU80 wild-type strain (p < 0.005). Only the Δgin4 strain exhibited a significant increase in apical compartment lengths (2.5-fold, p < 0.001; Figure 1C). Due to the observed similarity in the apical compartment length between the Δgin4 and ΔaspB strains, we next generated a Δgin4 ΔaspB double deletion strain to determine if Gin4 and AspB were acting in the same pathway or were contributing to septation through different mechanisms. The Δgin4 ΔaspB strain apical compartment length was similar to that of both the Δgin4 and ΔaspB single deletion strains, indicating that AspB and Gin4 may contribute to septation through the same pathway. In the basal compartments, the Δcla4 and ΔparA strains exhibited a hyperseptation phenotype, while the Δgin4 strain and the Δgin4 ΔaspB double deletion strain maintained the increased interseptal distances (Figure 1D).

Deletion of gin4 Results in Hypervirulence in an Invertebrate and a Murine Model of Invasive Aspergillosis

We previously reported that the ΔaspB strain was hypervirulent in the G. mellonella model but not in a murine model of invasive
FIGURE 2 | Deletion of parA increased susceptibility to Congo Red, but not to anti-cell wall drugs. Conidia (10⁴) from each strain were inoculated into GMM agar, GMM agar containing Congo Red (100 µg), Caspofungin (1 µg/ml), or Nikkomycin Z (2 µg/ml), and incubated for 3 days at 37°C.

aspergillosis. Similar to our findings with the ΔaspB strain, the Δgin4 strain resulted in a significantly increased mortality (100%) in G. mellonella by day +3 post infection, compared to the 70% mortality of the akuB<KU80> wild-type strain (p < 0.001; Figure 3A). In contrast, the ΔparA strain had a significantly reduced mortality in this model compared to the wild-type strain, and only 45% mortality at day +5 (p < 0.001). The Δcla4 strain displayed no statistical difference in mortality compared to the wild-type strain (p = 0.7971). Building on these invertebrate model virulence results, we then examined these strains in an intranasal murine model of invasive aspergillosis. The Δgin4 strain hypervirulence was reproduced in the murine model, with 100% mortality on day +6 (p < 0.001), while the Δcla4 and ΔparA strains were not significantly different from the wild-type strain (50–65% survival; p > 0.05; Figure 3B). To further understand the contribution of these proteins to pathogenesis, we performed histopathology analysis of the murine lungs after day +3 of infection (Figure 3C). Concomitant with the hypervirulence exhibited by the Δgin4 strain, there was a significant increase in fungal burden via Gomori’s methenamine silver staining in the wild-type akuB<KU80> and the Δgin4 strains. Although the Δcla4 and ΔparA strain mortality is similar to that of the wild-type strain, they differed in histopathology. The Δcla4 strain is defective in tissue invasion and only grew near the alveoli, while the ΔparA have a significant reduction in fungal burden. However, the Δcla4 and ΔparA strains infected mice presented symptoms of infection at the same level as the wild-type strain.
FIGURE 3 | The Δgin4 strain is hypervirulent in both an invertebrate host model and in a persistently immunosuppressed intranasal murine model of invasive aspergillosis. (A) Infection with the Δgin4 strain yield a 100% mortality by day +2 compared to 70% mortality in wild-type strain in the Galleria mellonella model (p < 0.001). The ΔparA strain was hypovirulent (60% survival by day +5) compared to the wild-type strain (p < 0.001). A total of 20 waxmoth G. mellonella larvae were infected with each A. fumigatus strain with 5 µl of a 1 × 10^8 spores/ml spore suspension. Infected larvae were incubated at 37 °C and survival was scored daily for 5 days. (B) Deletion of gin4 results in hypervirulence in a neutropenic murine model of invasive aspergillosis (p < 0.001). Mice were infected intranasally with a total of 4 × 10^6 conidia from each strain and survival monitored for 14 days. (C) Lung histopathology performed on day +3 days shows a significant increase in fungal burden (silver staining) and inflammation (H&E staining) between the wild-type and the Δgin4 strains.

AspB Exhibits Altered Localization Patterns in the Δgin4, Δcla4, and ΔparA Strains

During the multicellular growth stage, AspB localizes transiently at the septum, at possible septation sites (septal-like), as a ring, and in filaments and double bars (Figure 4) (Vargas-Muniz et al., 2015). Here, we found that AspB mislocalized in the Δgin4, Δcla4, and ΔparA backgrounds (Figure 4). In all the strains, including the wild-type strain, AspB localized as septal-like and double bars. In the ΔparA strain, the double bars had an intermediate length compared to the wild-type double bar localization. Although AspB in the wild type, Δgin4, and ΔparA strains localizes to ring-like structures, qualitative observation revealed that the rings in the Δgin4 and ΔparA background strains had a larger diameter compared to the rings observed in the wild-type strain. AspB localized into X-like structures in Δgin4 and ΔparA strains, similar to those observed in the wild-type strain after exposure to caspofungin. While AspB also localized in elongated X-like structures in the Δgin4 strain, it localized into intermediate length X-like structures in the ΔparA strain. In the Δgin4 and Δcla4 strains, AspB localized as a dot-like structure near the hyphal tip. Deletion of these kinases abolished the formation of AspB filaments, which remained present in the ΔparA strain. AspB in the Δgin4 strain localized into two different ring and bar structures, one that resembled the ring bar structures observed in the wild-type strain under caspofungin treatment, and another larger ring and bar that is unique to the Δgin4 strain. AspB localized at the majority of the hyphal tips in the Δcla4 strain, suggesting that the Cla4 kinase plays a role in excluding AspB from the hyphal tips. The structures described for each strain were commonly found throughout the mycelia. Compartments were observed to contain between 1-2 AspB structures, with the exception of multiple ring-like structures in the Δgin4 and ΔparA strains that were clumped together.

AspB Is Differentially Phosphorylated in the ΔparA Strain

In order to determine the role of Gin4, Cla4, and ParA as AspB posttranslational regulators, we first determined the phosphorylation profile of AspB in the wild-type strain. AspB
FIGURE 4 | Deletion of gin4, cla4, or para alters AspB localization pattern. Kinases Gin4 and Cla4, as well as the Protein Phosphatase 2A regulatory sub-unit ParA are required for proper localization of AspB under normal growth condition during the multicellular growth stage. The AspB–EGFP strain was grown in GMM broth for 20 h at 37°C and AspB–EGFP localization visualized by fluorescent microscopy. Scale bar, 2.5 µm.

was phosphorylated in seven residues: five in the GTPase domain (S134, S137, S247, T297, and T301), and two in the carboxyl-terminus after the septin unique element (S416 and S461) (Table 2 and Figure 5A). We next defined the effect of the deletions of gin4, cla4, and para on AspB phosphorylation in a comparative phosphoproteomic approach. This strategy revealed that only the para deletion altered the AspB phosphorylation profile, with AspB phosphorylated at two additional sites: T68 and S447 (Figures 5B–E). Quantitative proteomic (Skyline) analyses showed that both para specific phosphoresidues had a sevenfold increase in the area under the curve of the spectrogram compared to the wild-type strain (Table 3).

To assess the role of these phosphorylated residues as possible posttranslational regulation sites, we next generated two non-phosphorylatable as well as two phosphomimetic mutant strains (T68A and T68E; S447A and S447E). Phenotypic analyses of these phosphomutant strains revealed that locking AspB in either a phosphorylation-mimic or non-phosphorylable state resulted in mislocalization of AspB (Figure 6). Similar to the wild-type and the para strains, AspB phosphomutants localized into rings and double bars. However, the length of the bars observed was different in each of the AspB phosphomutants, while the ring-like structures were similar to those found in the para strain. Although all four AspB phosphomutants showed the localization of AspB to dot-like structures associated with the hyphal tip,
TABLE 2 | Phosphorylation sites within AspB identified by TiO₂ phospho-enrichment followed by LC–MS/MS analysis.

| Peptide sequence | Phosphorylated residue | Domain  | m/z          | Mascot ion score |
|------------------|------------------------|---------|--------------|------------------|
| TV[pS]ISADIEENGVR| S134                   | GTPase  | 82.16 + 0.1  | 22.4             |
| TVSIQ[pS]ISADIEENGVR| S137               | GTPase  | 87.63 + 0.09 | 25.2             |
| ADLTDE[pS]LFK   | S247                   | GTPase  | 82.88 + 0.08 | 24.3             |
| VPFAVGAN[pT]EVTADGR| T297               | GTPase  | 69.93 + 0.08 | 28.7             |
| VPFAVGANTE[pT]ADGR| T301               | GTPase  | 69.1 + 0.05  | 21.1             |
| LK[pS]EDEKYAR   | S416                   | C-terminus | 42.13 + 0.19 | 43.0             |
| KGF[pS]LR       | S461                   | C-terminus | 44.18 + 0.25 | 33.2             |

Aspergillus fumigatus Gin4 kinase is required for maintenance of proper interseptal distances and conidiation, similar to AspB. Deletion of aspB in the Δgin4 background exhibited a similar phenotype to the respective single deletion strains, suggesting that the Gin4 kinase and AspB may coordinately regulate septation. Nonetheless, AspB phosphorylation remains unchanged in the Δgin4 strain. This suggests that: (i) AspB might recruit Gin4 to possible septation sites and Gin4 then regulates septation, (ii) Gin4 indirectly regulates AspB through the phosphorylation of another yet unidentified AspB interacting partner, or (iii) Gin4 regulates AspB in a kinase-independent manner. Previous studies in S. cerevisiae have suggested that septins are required for proper localization and function of Gin4, and Gin4 phosphorylates Shs1 (non-core septin) and has functions that are independent from its kinase activity (Barral et al., 1998; Carroll et al., 1998; Longtine et al., 1998; Mortensen et al., 2002; Asano et al., 2006). Gin4 has been shown to regulate septins in a cell-cycle dependent manner in S. cerevisiae (Mortensen et al., 2002). It is still possible that Gin4 contributes to septin phosphorylation; however, this would be in a low abundant subset of AspB that we were not able to detect. Additionally, A. nidulans septins form distinct complexes at different growth stages (Hernandez-Rodiguez et al., 2014). The results reported here only focus on the multicellular growth stage, and it is possible we missed key regulatory phosphorylation events in the other growth stages. While the ΔaspB strain exhibited hypervirulence in the Galleria model but had no effect on virulence in the murine model (Vargas-Muniz et al., 2015), the Δgin4 strain is hypervirulent in both the insect and murine model of invasive aspergillosis. This could be due to misregulation of septin function in general or a septin-independent role of Gin4.

DISCUSSION

We previously demonstrated that the A. fumigatus core septin AspB is required for critical cellular processes, including regular septation, conidiation, and conidial cell wall organization (Vargas-Muniz et al., 2015); however, regulation of AspB remains unclear. In S. cerevisiae, phosphorylation has been described as a possible mechanism of septin regulation (Tang and Reed, 2002; Garcia et al., 2011). Here, we explored the role of Gin4 and Cla4 kinases and the regulatory subunit of protein phosphatase 2A, ParA, as possible regulators of AspB. We found that AspB is phosphorylated at seven residues in vivo at the multicellular growth stage. Previously, core septins have been reported to be generally phosphorylated in the divergent amino- and carboxyl-termini, with the exception of S. cerevisiae Cdc10 (Hernandez-Rodriguez and Momany, 2012). However, most of the A. fumigatus AspB phosphorylation sites are within the GTPase domain. This is interesting because the majority of the known phosphorylation sites in septins are not present in the GTPase domain. While these residues are conserved across the filamentous ascomycetes, the same is not the case in other species. Due to the scattered pattern of septin phosphorylations observed in other species we expect no conservation in the residues phosphorylated (Hernandez-Rodriguez and Momany, 2012). However, further phosphoproteomic characterization is required to determine if phosphorylation of these residues is a conserved mechanism in other filamentous ascomycetes.
FIGURE 5 | AspB is differentially phosphorylated in the ΔparA strain. (A) Schematic representation of AspB conserved domains and phosphorylation sites. Black bars indicate AspB phosphorylation sites in the wild-type strain and red bars indicate phosphorylation sites unique to the ΔparA strains. (B–E) AspB–EGFP and ΔparA AspB–EGFP strains were grown in GMM liquid for 24 h at 37°C. AspB–EGFP was purified using GFP-Trap® affinity purification, digested with trypsin and phospho-enriched using TiO<sub>2</sub>. LC–MS/MS was used to identify phosphopeptides. X-axis represents intensity of the signal, and Y-axis represents the retention time of the phosphopeptide in minutes after mixture injection to the liquid chromatography column. (B,C) Chromatogram for the KLTGYVGFANLPNQWHR peptide in (B) wild-type and (C) ΔparA strains. (C,D) LESGRPIEEK phosphopeptides chromatogram in (D) wild-type and (E) ΔparA strains. Area under the curve from each chromatogram peak was used for the Skyline analysis.

but the hyphal compartment was significantly reduced (Li et al., 2004). In *C. albicans*, deletion of *cla4* resulted in reduction in fungal burden and hyphal invasion in mouse kidneys, leading to a reduction in pathogenesis (Leberer et al., 1997). The *A. fumigatus* Δ*cla4* strain also showed decreased tissue invasion in the intranasal murine model of invasive aspergillosis, however; the Δ*cla4* strain remained as virulent as the *akuB* KU80 strain. The Δ*cla4* strain did not phenocopy the Δ*aspB* strain or alter the

| Peptide sequence | Phosphorylated residue | m/z    | *akuB* KU80 Area | Δ parA Area | Fold change |
|------------------|------------------------|--------|-----------------|-------------|-------------|
| KLTPTG/Y/GFA/NP/QWHR | T68                    | 76.06 ± 0.05 | 46987743       | 327294944   | 6.97        |
| LEpS[GRPIEEK     | S447                   | 32.84 ± 0.12  | 55234844       | 8304621.5   | 6.65        |
AspB phosphorylation profile. Nonetheless, the AspB localization pattern in the Δcla4 strain was altered. It is suggested that Cla4 is capable of regulating the actin cytoskeleton in some fungi, and Cla4 could have a similar role in A. fumigatus (Leveleki et al., 2004). Furthermore, previously we showed that septin localization is dependent on actin and microtubules, and one of the septins, AspE, interacts with actin and microtubules (Juvvadi et al., 2011b, 2013). Thus, this altered localization could be a result of mislocalization of actin due to the absence of the Cla4 kinase.

Similar to A. nidulans, A. fumigatus ParA is involved in hyphal growth, conidiation and septation (Zhong et al., 2014). The ΔparA strain was hypovirulent in the G. mellonella model; however, the survival and histopathological analyses of the ΔparA strain were similar to that of the wild-type strain in the murine model. This difference in virulence between the two models could be due to the immunosuppression used in the murine model to establish infection, compared to the immunocompetent invertebrate model with a rudimentary immune system. Therefore, ParA is dispensable for pathogenesis in an immunosuppressed mammalian host. The role of RTS1 (ParA ortholog) as a septin regulator has been previously explored in S. cerevisiae, where it coordinates the dynamics of the septin rings by Rts1-dependent dephosphorylation of Shs1 (Dobbelaere et al., 2003). In A. fumigatus, deletion of parA resulted in altered AspB localization and, most importantly, altered the phosphorylation profile of AspB revealing ParA-dependent dephosphorylation of AspB. AspB under normal growth conditions is phosphorylated in seven residues distributed across the GTPase domain and the C-terminal region. Deletion of parA resulted in two additional phosphorylation sites (T68 and S447) that are enriched sevenfold when compared to the control strain. Our preliminary studies on the AspB interactome using purification of the AspB–EGFP fusion protein using GFP-Trap affinity matrix and subsequent LC–MS/MS analysis revealed PP2A as one of the AspB interactants, suggesting that PP2A could be directly

![AspB localization patterns in different AspB phosphomutant strains.](image-url)
dephosphorylating these residues (data not shown). Although mutation of these sites (S447A, S447E, T68A, and T68E) altered the localization pattern of AspB, only apical compartment length was affected in the AspB^{S447A}{T68E}. Interestingly, AspB^{S447A}{T68E} localizes transiently at the septum, similar to wild-type AspB, and the defect in apical compartment length could be due to altered interaction with septation regulators due to locking AspB in a phosphomimetic state. T68 is in the analogous position to A. gossypii S91 that has also been reported to be phosphorylated in vivo (Meseroll et al., 2013). A phosphomimetic strain of S91 in A. gossypii also resulted in a modest phenotype with abnormal spore morphology; however, spores were viable and capable of germinating at the same rate as the wild type. More detailed characterization of these mutants, as well as phosphomutants of the other AspB phosphoiresidues, will further our understanding on how AspB is regulated through phosphorylation and how these phosphorylation sites might affect AspB–protein interactions.

**AUTHOR CONTRIBUTIONS**

JV and WS conceived and designed research; JV, HR, and AR performed research; JV, HR, AR, and YA performed virulence and histopathology analyses; JV GW, ES, and MM acquired and analyzed the phosphoproteomic data. JV, PJ, and WJ wrote the paper; JV, HR, AR, GW, ES, MM, YA, PJ, and WS approved the final submission.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: http://journal.frontiersin.org/article/10.3389/fmicb.2016.00997

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